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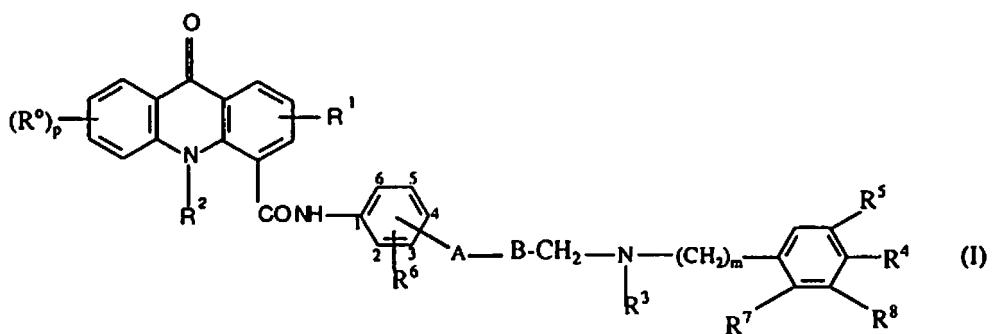
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㉓ Acridine derivatives.

㉔ There are provided compounds of the general formula (I)



wherein R⁰ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio, amino or nitro group;

p represents 1; or when R⁰ represents C₁₋₄alkoxy may also represent 2 or 3;

R¹ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy or C₁₋₄alkylthio group;

R² represents a hydrogen atom or a C₁₋₄alkyl group;

A represents an oxygen or a sulphur atom, a bond or a group (CH₂)₁NR³ (where 1 represents zero or 1 and R³ represents a hydrogen atom or a methyl group);

B represents a C₁₋₄alkylene chain optionally substituted by a hydroxyl group, except that the hydroxyl group and moiety A cannot be attached to the same carbon atom when A represents an oxygen or sulphur atom or a group (CH₂)₁NR³, or when A represents a bond B may also represent a C₂₋₄alkenylene chain;

R³ represents a hydrogen atom or a C₁₋₄alkyl group;

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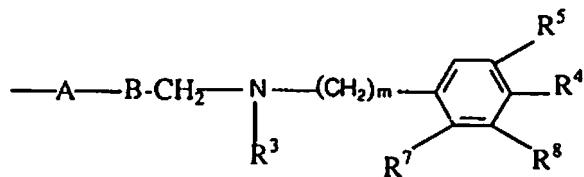
m represents 1 or 2;

R^4 represents a hydrogen or a halogen atom, or a C_1 - 4 alkyl, C_1 - 4 alkoxy or C_1 - 4 alkylthio group;

R^5 represents a hydrogen atom or a C_1 - 4 alkoxy group;

R^6 represents a hydrogen atom or a C_1 - 4 alkyl or C_1 - 4 alkoxy group R^7 represents a hydrogen atom or R^3 and R^7 together form a group $-(CH_2)_n$ -where n represents 1 or 2;

R^8 represents a hydrogen atom or a C_1 - 4 alkoxy group;
the group



is attached at the benzene ring 3 or 4 position relate to the carboxamide substituent, provided that when the group is attached at the benzene ring 3 position then R^6 must be attached at the benzene ring 6 position;
and salts and solvates thereof.

The novel compounds of formula (I) can sensitize multidrug-resistant cancer cells to chemotherapeutic agents and may be formulated for use in therapy, particularly to improve or increase the efficacy of an antitumour drug.

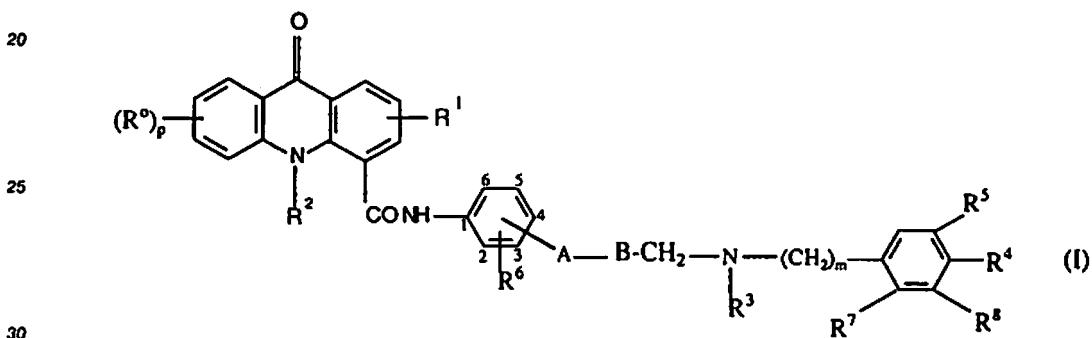
This invention relates to acridine derivatives, to processes for their preparation, to pharmaceutical compositions containing them, and to their medical use. In particular it relates to compounds and compositions which are capable of sensitizing multidrug-resistant cancer cells to chemotherapeutic agents.

In many patients, the efficacy of cancer chemotherapy is initially poor or decreases after initial treatment due to the development of resistance to anticancer drugs, known as multidrug-resistance. Multidrug-resistance is a process whereby malignant cells become resistant to structurally diverse chemotherapeutic agents following treatment with a single anti-tumour drug. This acquired drug resistance can be a major clinical obstacle in the treatment of cancer. Some tumours are intrinsically multidrug-resistant, and hence do not respond to chemotherapy.

It has been shown that this type of resistance can be reversed by some calcium channel blockers such as nifedipine and verapamil, by antiarrhythmic agents such as amiodarone and quinidine, as well as by natural products such as cephalexin. However, these compounds exert their multidrug resistant cell sensitizing activity only at very high doses, well above their intrinsic toxic level, and this severely limits their clinical use in the field of cancer chemotherapy.

A novel group of compounds has now been found which can sensitize multidrug-resistant cancer cells to chemotherapeutic agents at dose levels at which these novel compounds show no toxicity.

Thus, the present invention provides a compound of formula (I):



wherein R⁰ represents a hydrogen or halogen atom, or a C₁-4 alkyl, C₁-4 alkoxy, C₁-4 alkylthio, amino or nitro group;

p represents 1; or when R⁰ represents C₁-4 alkoxy may also represent 2 or 3;

35 R¹ represents a hydrogen or halogen atom, or a C₁-4 alkyl, C₁-4 alkoxy or C₁-4 alkylthio group;

R² represents a hydrogen atom or a C₁-4 alkyl group;

A represents an oxygen or a sulphur atom, a bond or a group (CH₂)₁NR³ (where 1 represents zero or 1 and R³ represents a hydrogen atom or a methyl group);

40 B represents a C₁-4 alkylene chain optionally substituted by a hydroxyl group, except that the hydroxyl group and moiety A cannot be attached to the same carbon atom when A represents an oxygen or sulphur atom or a group (CH₂)₁NR³, or when A represents a bond B may also represent a C₂-4 alkenylene chain;

R³ represents a hydrogen atom or a C₁-4 alkyl group;

m represents 1 or 2;

R⁴ represents a hydrogen or a halogen atom, or a C₁-4 alkyl, C₁-4 alkoxy or C₁-4 alkylthio group;

45 R⁵ represents a hydrogen atom or a C₁-4 alkoxy group;

R⁶ represents a hydrogen atom or a C₁-4 alkyl or C₁-4 alkoxy group;

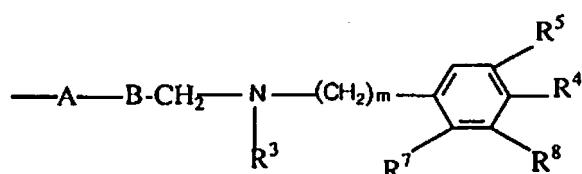
R⁷ represents a hydrogen atom or R³ and R⁷ together form a group -(CH₂)_n- where n represents 1 or 2;

R⁸ represents a hydrogen atom or a C₁-4 alkoxy group;

the group

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is attached at the benzene ring 3 or 4 position relate to the carboxamide substituent, provided that when the group is attached at the benzene ring 3 position then R⁶ must be attached at the benzene ring 6 position; and salts and solvates thereof including physiologically acceptable salts and solvates thereof.

As used herein, an alkyl group, either as such or as part of an alkoxy or alkylthio group may be a straight chain or branched chain alkyl group, for example a methyl, ethyl or prop-2-yl group.

A halogen substituent may be a fluorine, chlorine, bromine or iodine atom.

The group(s) R⁰, when other than a hydrogen atom, may be situated at the 5, 6, 7 or 8-position of the acridone molecule, and the group R¹, when other than a hydrogen atom, may be situated at the 1, 2 or 3-position of the acridone molecule.

Examples of the chain -A-B-CH₂- include -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, -CH₂NMe(CH₂)₂-, -CH=CHCH₂-, -CH₂CH=CHCH₂-, -CH(OH)CH₂-, -O(CH₂)₂-, -O(CH₂)₃-, -OCH₂CH(OH)CH₂-, -NH(CH₂)₂-, -S(CH₂)₂- and -S(CH₂)₃-.

A preferred class of compounds of formula (I) is that in which R⁰ represents a hydrogen or fluorine atom, or a C₁₋₄alkoxy (e.g. methoxy) group, C₁₋₄alkyl (e.g. methyl) or C₁₋₄alkylthio (e.g. methylthio) group, and R¹ is a hydrogen atom. When R⁰ represents a substituent other than a hydrogen atom, an R⁰ group is preferably situated at the 5-position of the acridone molecule.

Another preferred class of compounds of formula (I) is that in which R² represents a hydrogen atom.

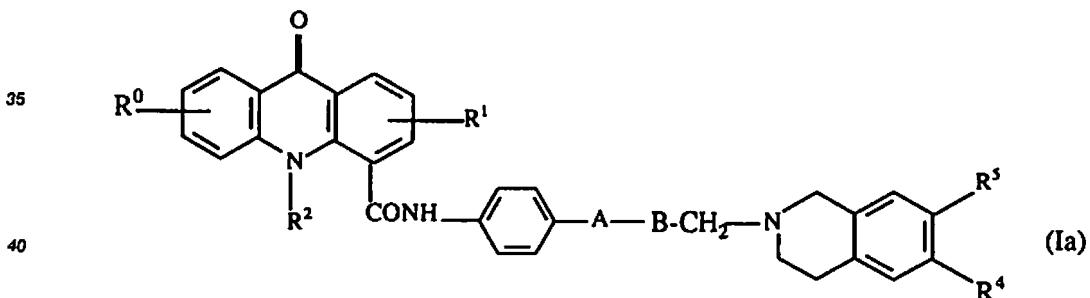
When R³ represents a hydrogen atom or a C₁₋₄alkyl group, preferably R³ represents a C₁₋₄alkyl (e.g. methyl) group.

Yet another preferred class of compounds of formula (I) is that in which R⁴ represents a hydrogen atom or a C₁₋₄alkoxy (e.g. methoxy) group, R⁵ represents a hydrogen atom or a C₁₋₄alkoxy (e.g. methoxy) group and R⁶ represents a hydrogen atom or a C₁₋₄alkoxy (e.g. methoxy) group, provided that at least one of R⁴, R⁵ and R⁶ represents a C₁₋₄alkoxy (e.g. methoxy) group. A particularly preferred class of compounds of formula (I) is that in which R⁴ and R⁵ each represent a C₁₋₄alkoxy (e.g. methoxy) group and R⁶ represents a hydrogen atom.

A further preferred class of compounds of formula (I) is that in which R⁶ represents a hydrogen atom or a methyl, ethyl, methoxy or ethoxy group.

A preferred group of compounds of formula (I) is that in which m represents 1 and R³ and R⁷ together form a group -(CH₂)₂-, and physiologically acceptable salts and solvates thereof.

A particular group of compounds of formula (I) is that of formula (Ia)



wherein R⁰ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio or nitro group;

R¹ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy or C₁₋₄alkylthio group;

R² represents a hydrogen atom or a C₁₋₄alkyl group;

A represents an oxygen or a sulphur atom or a bond;

B represents an unsubstituted C₁₋₄alkylene chain;

50 R⁴ and R⁵ each independently represents a C₁₋₄alkoxy group; and physiologically acceptable salts and solvates thereof.

A particularly preferred group of compounds of formula (I) is that of formula (Ia) in which R⁰ represents a hydrogen or fluorine atom or a C₁₋₄alkoxy (e.g. methoxy) or C₁₋₄alkyl (e.g. methyl) group, R¹ and R² each represent a hydrogen atom and R⁴ and R⁵ each represent a C₁₋₄alkoxy (e.g. methoxy) group. Such compounds in which the R⁰ group is situated at the 5-position of the acridone molecule are especially preferred.

It is to be understood that the present invention includes all combinations of the aforementioned particular and preferred groups.

A particularly preferred compound according to the invention is 9,10-dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide and physiologically acceptable salts and solvates thereof.

Other preferred compounds according to the invention are:-

- 5 9,10-dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide;
- 5-fluoro-9,10-dihydro-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide;
- 9,10-dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-4-acridinecarboxamide;
- 10 9,10-dihydro-5-methyl-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide;
- 9,10-dihydro-5-methoxy-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-9-oxo-4-acridinecarboxamide;
- 15 9,10-dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-5-methyl-9-oxo-4-acridinecarboxamide;
- and physiologically acceptable salts and solvates thereof.

Further preferred compounds according to the invention are :-

- N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- 20 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- 25 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
- and physiologically acceptable salts and solvates thereof.

Yet further preferred compounds according to the invention are:-

- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- 30 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
- 35 and physiologically acceptable salts and solvates thereof.

Other preferred compounds according to the invention are :-

- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
- 40 N-[4-[3-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
- 45 N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[2-[(3,4-dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-
- 50 acridinecarboxamide;
- N-[4-[5-[(3,4-dimethoxyphenyl)methyl]methylamino]pentyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
- 55 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethylamino] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
- N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-fluoro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methylthio-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide;

5 N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide;
 and physiologically acceptable salts and solvates thereof.

Yet further preferred compounds according to the invention are :-

N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

10 N-[4-[4-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(2-(4-methoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-2-(methylthio)-9-oxo-4-acridinecarboxamide;

15 N-[4-[3-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(2-(4-methoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(3,4-dimethoxyphenyl)methyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

20 N-[4-[3-[(3,4-dimethoxyphenyl)methyl)methylamino]propoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(3,4-dimethoxyphenyl)methyl)methylamino]ethyl]thio] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;

25 and physiologically acceptable salts and solvates thereof.

Suitable physiologically acceptable salts of the compounds of formula (I) include acid addition salts formed with organic or inorganic acids, for example, hydrochlorides, hydrobromides, sulphates, alkyl- or arylsulphonates (e.g. methanesulphonates or *p*-toluenesulphonates), phosphates, acetates, citrates, succinates, lactates, tartrates, fumarates and maleates. The solvates may, for example, be hydrates.

30 Other salts which are not physiologically acceptable may be useful in the preparation of compounds of formula (I) and these form a further part of the invention.

The ability of the compounds of formula (I) to sensitize multidrug-resistant cells has been demonstrated *in vitro* in the multidrug-resistant Chinese hamster ovary cell line (described by Bech-Hansen et al., *J Cell. Physiol.*, 1976, 88, 23-32) and the multidrug-resistant human mammary carcinoma line (described by Batist et al., (*J. Biol. Chem.*, 1986, 261, 1544-1549) using an assay similar to that described by Carmichael et al., *Cancer Research*, 1987, 47, 936.

The ability of the compounds of formula (I) to sensitize multidrug-resistant cells has also been demonstrated *in vivo* in the tumour line P388R (described by Johnson et al., *Cancer Treat. Rep.*, 1978, 62, 1535-1547). The methodology used is similar to that described by Boesch et al., *Cancer Research*, 1991, 51, 4226-4233. However, in our study the compounds were administered orally, intravenously or intraperitoneally in a single dose.

The present invention accordingly provides a compound of formula (I) or a physiologically acceptable salt or solvate thereof for use in therapy, more particularly for use in the treatment of a mammal, including a human, which is suffering from cancer to:

45 (a) improve or increase the efficacy of an antitumour drug; or
 (b) increase or restore sensitivity of a tumour to an antitumour drug; or
 (c) reverse or reduce resistance, whether acquired, induced or innate, of a tumour to an antitumour drug.

The present invention also provides a method of treatment of a mammal, including a human, which is suffering from cancer, which method comprises administering to said mammal an effective amount of a compound of formula (I) or a physiologically acceptable salt or solvate thereof to:

(a) improve or increase the efficacy of an antitumour drug; or
 (b) increase or restore sensitivity of a tumour to an antitumour drug; or
 (c) reverse or reduce resistance, whether acquired, induced or innate, of a tumour to an antitumour drug.

In another aspect, the present invention provides the use of a compound of formula (I) or a physiologically acceptable salt or solvate thereof for the manufacture of a medicament for the treatment of a mammal, including a human, which is suffering from cancer to:

(a) improve or increase the efficacy of an antitumour drug; or
 (b) increase or restore sensitivity of a tumour to an antitumour drug; or

(c) reverse or reduce resistance, whether acquired, induced or inate, of a tumour to an antitumour drug.

It will be appreciated that the compounds according to the present invention are administered in conjunction with an antitumour drug. Thus, in a further aspect, the present invention provides a product containing a compound of formula (I) or a physiologically acceptable salt or solvate thereof and an antitumour drug as a combined preparation for simultaneous, separate or sequential use in treating cancer, more particularly to:

- (a) improve or increase the efficacy of said antitumour drug; or
- (b) increase or restore sensitivity of a tumour to an antitumour drug; or
- (c) reverse or reduce resistance, whether acquired, induced or inate, of a tumour to an antitumour drug.

Examples of suitable antitumour drugs for use in conjunction with compounds of the present invention include Vinca alkaloids (e.g. vincristine, vinblastine and vinorelbine), anthracyclines (e.g. daunorubicin, doxorubicin and aclarubicin), taxol and derivatives thereof (e.g. taxotere), podophyllotoxins (e.g. etoposide and VP16), mitoxantrone, actinomycin, colchicine, gramicidine D, amsacrine or any drug having cross-resistance with the above drugs characterised by the so-called MDR phenotype.

It will be appreciated that if administration of the two drugs is not simultaneous, the delay in administering the second of the active ingredients should not be such as to lose the beneficial effect of the combination.

Thus, in a further aspect, the present invention provides a compound of formula (I) or a physiologically acceptable salt or solvate thereof and an anticancer drug in the presence of each other in the human or non-human animal body for use in treating cancer, more particularly to:

- (a) improve or increase the efficacy of said antitumour drug; or
- (b) increase or restore sensitivity of a tumour to an antitumour drug; or
- (c) reverse or reduce resistance, whether acquired, induced or inate, of a tumour to an antitumour drug.

Some tumours are often intrinsically multidrug-resistant, notably colon carcinomas, renal cell carcinomas, hepatomas and adrenocortical carcinomas.

Other types of tumour are often initially sensitive but can become multidrug-resistant, notably leukaemias, lymphomas, myelomas, paediatric tumours (e.g. neuroblastomas), sarcomas, and breast, ovarian and lung cancers.

Hence the compounds of the invention are particularly useful in the treatment of mammals, including humans, receiving chemotherapy for one of the above types of cancer.

In using a compound of formula (I) or a physiologically acceptable salt or solvate thereof and an antitumour drug it may be preferable to employ the active ingredients in the form of separate pharmaceutical formulations, although a single combined formulation can be used as demonstrated hereinafter. However, in the latter formulation both active ingredients must of course be stable and mutually compatible in the particular formulation employed.

Pharmaceutical formulations of suitable antitumour drugs and appropriate dosages and dosage rates will generally correspond with those one would use if administering the antitumour drug alone to treat a tumour.

Suitable pharmaceutical formulations and appropriate dosages and dosage rates of compounds of formula (I) and physiologically acceptable salts and solvates thereof are described hereinafter.

Thus, in a further aspect, the invention provides a pharmaceutical composition which comprises a compound of formula (I) or a physiologically acceptable salt or solvate thereof together with one or more physiologically acceptable carriers or excipients.

In another aspect, the present invention provides a pharmaceutical composition which comprises an active amount of a compound of formula (I) or a physiologically acceptable salt or solvate thereof for use in the treatment of a mammal which is suffering from cancer, to:

- (a) improve or increase the efficacy of an antitumour drug; or
- (b) increase or restore sensitivity of a tumour to an antitumour drug; or
- (c) reverse or reduce resistance, whether acquired, induced or inate, of a tumour to an antitumour drug.

The compounds according to the invention may be formulated for oral, buccal, parenteral or rectal administration, of which oral and parenteral are preferred.

For oral administration, the pharmaceutical compositions may take the form of, for example, tablets or capsules prepared by conventional means with pharmaceutically acceptable excipients such as binding agents (e.g. pre-gelatinised maize starch, polyvinylpyrrolidone or hydroxypropyl methylcellulose); fillers (e.g. lactose, microcrystalline cellulose or calcium hydrogen phosphate); lubricants (e.g. magnesium stearate, talc or silica); disintegrants (e.g. sodium lauryl sulphate or sodium starch glycolate). The tablets may be coated by methods well known in the art. Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may be prepared by conventional

means with pharmaceutically acceptable additives such as suspending agents (e.g. sorbitol syrup, cellulose derivatives or hydrogenated edible fats); emulsifying agents (e.g. lecithin or acacia); non-aqueous vehicles (e.g. almond oil, oily esters, ethyl alcohol or fractionated vegetable oils); and preservatives (e.g. methyl or propyl-p-hydroxybenzoates or sorbic acid). The preparations may also contain buffer salts, flavouring, 5 colouring and sweetening agents as appropriate.

Preparations for oral administration may be suitably formulated to give controlled release of the active compound.

For buccal administration the compositions may take the form of tablets or lozenges formulated in conventional manner.

10 The compounds of the invention may be formulated for parenteral administration by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form e.g. in ampoules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily, aqueous or alcoholic vehicles, and may contain formulatory agents such as suspending, stabilising and/or dispersing agents. Alternatively, the active ingredient may be in powder form 15 for constitution with a suitable vehicle, e.g. sterile pyrogen-free water, before use.

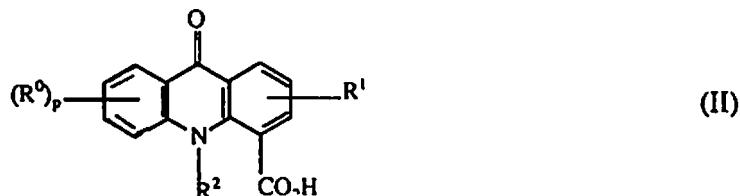
The compounds of the invention may also be formulated in rectal compositions such as suppositories or retention enemas, e.g. containing conventional suppository bases such as cocoa butter or other glycerides.

20 A proposed daily dose of the compounds of the invention for administration to a human (of approximately 70kg body weight) is about 10mg to 1000mg, more preferably about 25mg to 500mg. It will be appreciated that it may be necessary to make routine variations to the dosage, depending on the age and 25 condition of the patient, and the route of administration. For example, a daily dose of about 1mg/kg may be appropriate for administration to a human by infusion. The daily dose may be given as a single unit or as two or more subunits administered after appropriate time intervals.

25 Compounds of general formula (I) and physiologically acceptable salts and solvates thereof may be prepared by the general methods outlined hereinafter. In the following description, the groups R⁰ to R⁸, m, p, A and B are as defined for compounds of formula (I) unless otherwise specified.

Thus according to a first general process (A), a compound of formula (I) may be prepared by reacting a compound of formula (II):

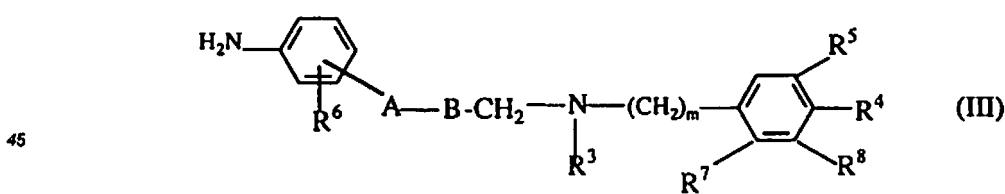
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with a compound of formula (III)

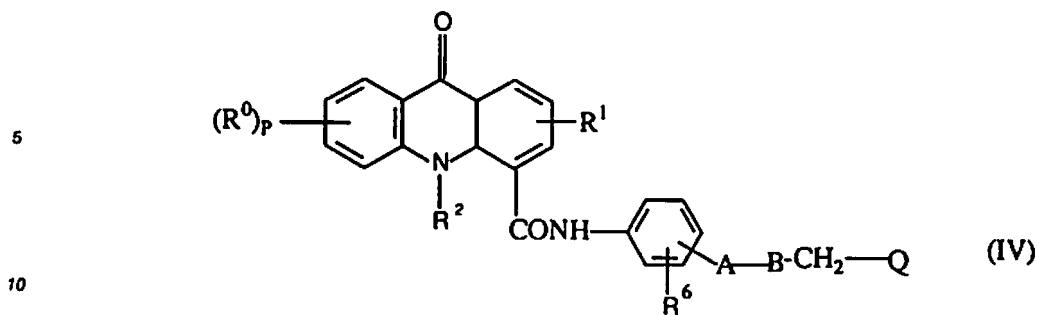
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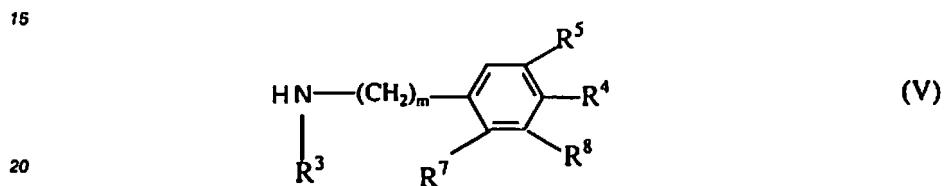
45

50 The reaction may be effected using a coupling reagent standardly used in peptide synthesis, such as dicyclohexylcarbodiimide (optionally in the presence of 1-hydroxybenzotriazole), diphenylphosphoryl azide or N,N'- carbonyldiimidazole. The reaction may be conveniently effected in an inert solvent such as an ether (e.g. tetrahydrofuran), a halogenated hydrocarbon (e.g. dichloromethane), an amide (e.g. dimethylformamide) or a ketone (e.g. acetone), and at a temperature of, for example, -10 to + 100°C, more preferably at about room temperature.

55 According to another general process (B), a compound of formula (I) may be prepared by reacting a compound of formula (IV):

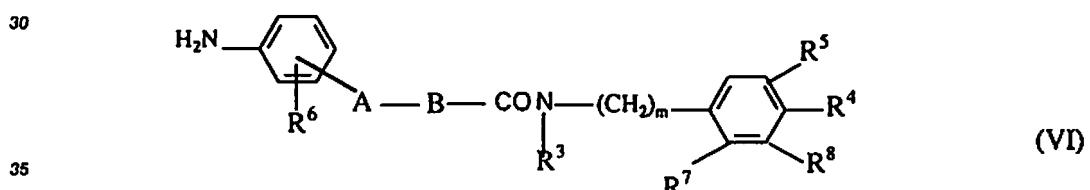


wherein Q represents a halogen (e.g. bromine) atom, with a compound of formula (V):



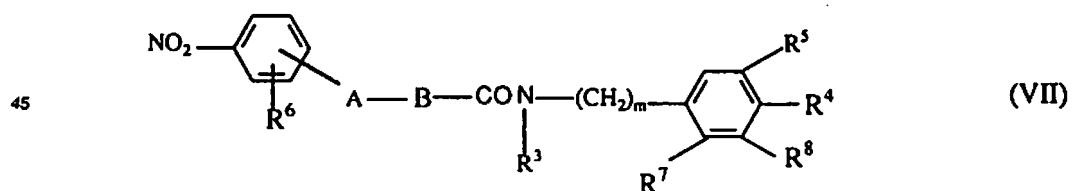
or a salt thereof. The reaction may be effected in the presence of an acid acceptor such as an alkali metal carbonate (e.g. potassium carbonate), in the presence or absence of a solvent, at an elevated temperature (e.g. 50 to 120 °C). Suitable solvents include ketones (e.g. acetone, methylethylketone or methylisopropylketone) and alcohols (e.g. ethanol or isopropanol).

Compounds of formula (III) in which A represents an oxygen atom or a bond may be prepared by the reduction of a compound of formula (VI):



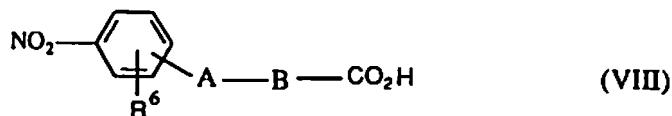
(in which A is an oxygen atom or a bond) with a suitable reducing agent such as lithium aluminium hydride in an inert solvent such as an ether (e.g. tetrahydrofuran) at an elevated temperature.

40 Compounds of formula (VI) may be prepared by the reduction of a compound of formula (VII):



50 by catalytic hydrogenation, for example using hydrogen in the presence of a noble metal catalyst (e.g. palladium). The catalyst may be supported on, for example, charcoal. The hydrogenation may be effected in a solvent such as an alcohol (e.g. ethanol), and conveniently at a temperature in the range of 20° to 100° C (e.g. 20° to 50° C) and atmospheric pressure. Alternatively, the reduction may be effected using iron and concentrated hydrochloric acid at an elevated temperature (e.g. reflux). This alternative reduction procedure leaves any double bond present in the compound of formula (VII) intact.

55 Compounds of formula (VII) may be prepared by the reaction of a compound of formula (VIII):

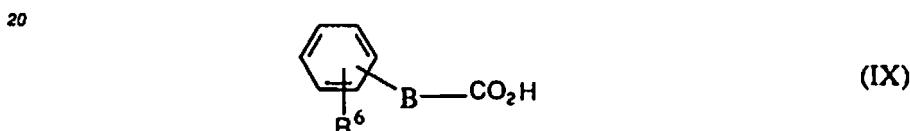


or an activated derivative thereof, with a compound of formula (V) as defined previously or a salt thereof, optionally in the presence of a base such as an organic base (e.g. triethylamine or *N,N*-diisopropylethylamine) or an inorganic base such as an alkali metal carbonate (e.g. potassium carbonate) or 10 hydrogen carbonate (e.g. sodium hydrogen carbonate).

When the free acid (VIII) is reacted with the amine (V), coupling reagents and conditions described in process (A) for the reaction of a compound of formula (III) with a compound of formula (III) may be used.

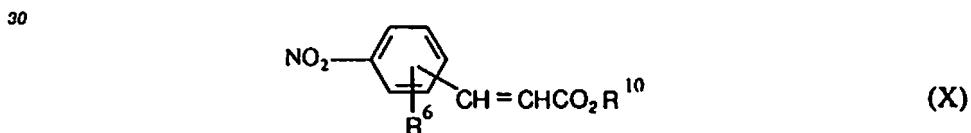
When an activated derivative of a compound of formula (VIII) is used, this may be, for example, an acid halide (e.g. an acid chloride), prepared by reacting the free acid (VIII) with a halogenating reagent (e.g. 15 thionyl chloride). This activated derivative of a compound of formula (VIII) may be reacted with a compound of formula (V) in a solvent such as acetone in the presence of a base such as sodium hydrogen carbonate.

Compounds of formula (VIII) wherein A represents a bond may be prepared by the nitration of a compound of formula (IX):



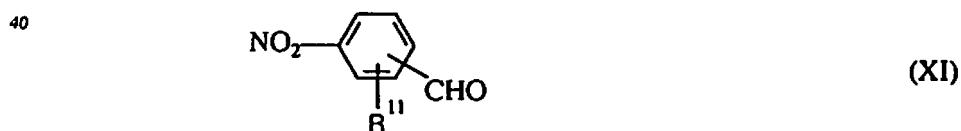
25 with nitric acid.

Compounds of formula (VIII) wherein A represents a bond and B represents a group -CH=CH- may conveniently be prepared by the hydrolysis of a compound of formula (X):



35 where R¹⁰ represents a C₁₋₄alkyl group. The hydrolysis may be effected using conventional methods, for example, by using sodium hydroxide in aqueous ethanol.

Compounds of formula (X) may be prepared by the reaction of a compound of formula (XI):

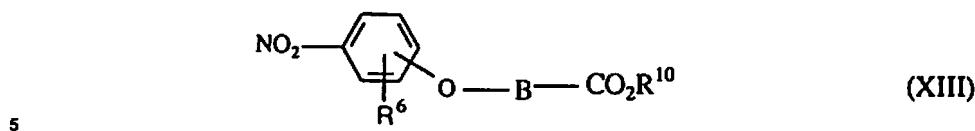


45 where R¹¹ represents a hydrogen atom or a C₁₋₄alkyl, C₁₋₄alkoxy or hydroxyl group, with a compound of formula (XII):



where R¹⁰ is as defined previously, in an inert solvent such as a hydrocarbon (e.g. toluene) and at an elevated temperature. For the preparation of a compound of formula (X) wherein R⁶ represents a C₁₋₄alkoxy group from a compound of formula (XI) wherein R¹¹ represents a hydroxyl group, the above reaction is followed by alkylation of the hydroxyl group using, for example, an alkyl halide.

55 Compounds of formula (VIII) wherein A represents an oxygen atom may be prepared by the hydrolysis of a compound of formula (XIII):



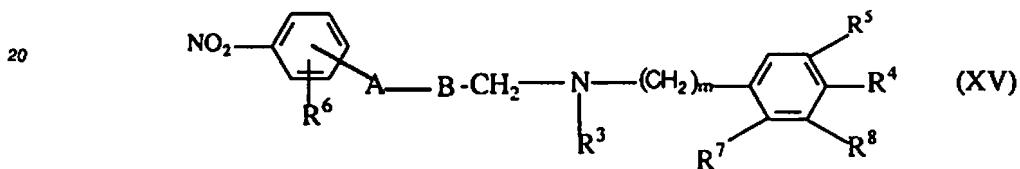
wherein R^{10} is as defined above. The hydrolysis may be effected using conventional methods, for example, by using sodium hydroxide in aqueous ethanol.

Compounds of formula (XIII) may be prepared by the reaction of a compound of formula (XIV):



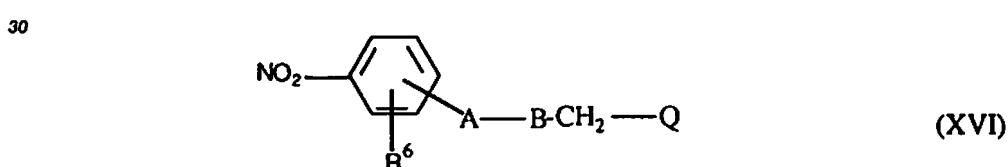
wherein L represents a halogen (e.g. bromine) atom, with a nitrophenol derivative in the presence of an alkali metal carbonate (e.g. potassium carbonate), in a solvent such as acetone.

15 Compounds of formula (III) wherein A represents an oxygen or sulphur atom or a bond may also be prepared by the reduction of a compound of formula (XV):



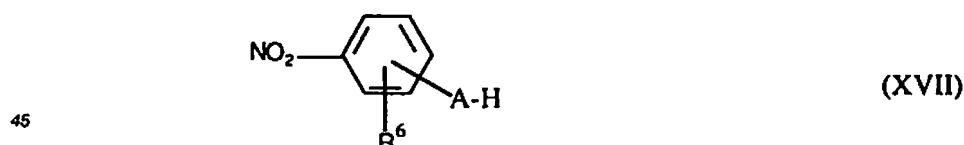
25 (where A is an oxygen or sulphur atom or a bond) using the conditions described above for the reduction of a compound of formula (VII).

Compounds of formula (XV) may be prepared by heating a compound of formula (XVI):



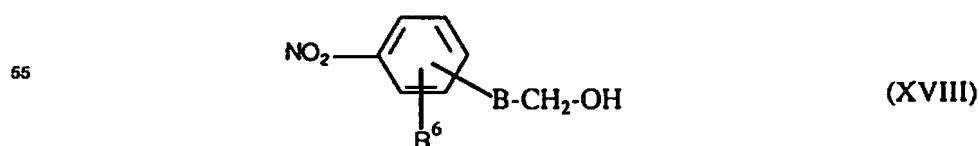
(wherein Q represents a halogen (e.g. bromine) atom and A is an oxygen or sulphur atom or a bond), with a compound of formula (V) as defined above under the conditions described in process (B) above.

40 Compounds of formula (XVI) wherein A represents an oxygen or a sulphur atom may be prepared by the reaction of a compound of formula (XVII):



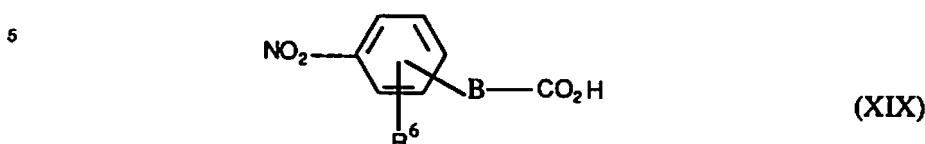
wherein A represents an oxygen or a sulphur atom, with a dihaloalkane Q-B-CH2-Q in the presence of a suitable base such as an alkali metal carbonate (e.g. potassium carbonate).

50 Compounds of formula (XVI) wherein A represents a bond may be prepared by the reaction of a compound of formula (XVIII):



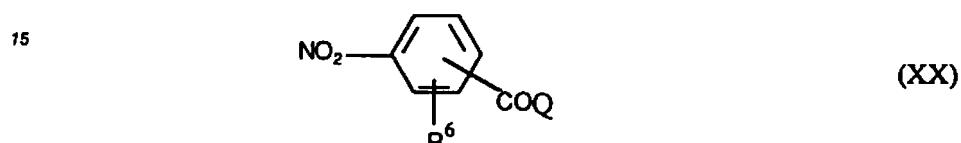
with an halogenating reagent such as phosphorus tribromide.

Compounds of formula (XVIII) may be prepared by the reduction of a compound of formula (XIX):



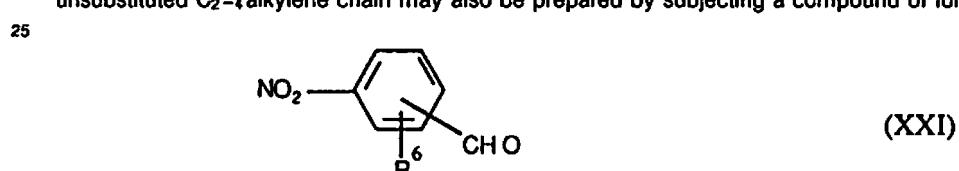
10 with a suitable reducing agent such as diborane.

Compounds of formula (XIX) may be prepared by subjecting a compound of formula (XX):



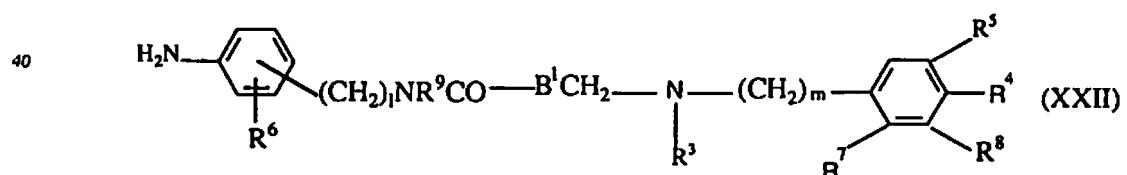
20 wherein Q represents a halogen (e.g. chlorine) atom to one or more successive Arndt-Eistert syntheses (i.e. reaction with diazomethane followed by treatment with, for example, silver oxide and water).

It will be appreciated by one skilled in the art that compounds of formula (XIX) in which B represents an unsubstituted C₂-4 alkylene chain may also be prepared by subjecting a compound of formula (XXI):



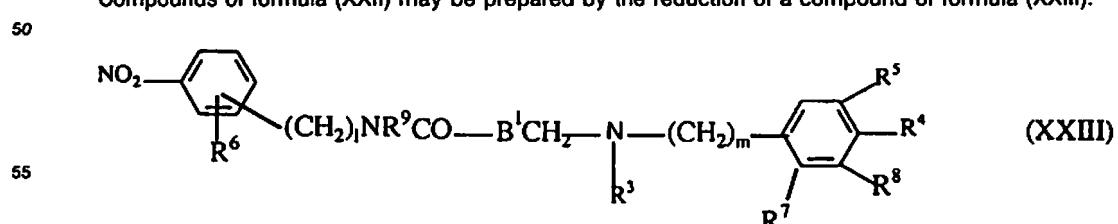
30 to a Wittig reaction with a suitable phosphorus ylid (e.g. Ph₃P = CH(CH₂)₃OH) followed by reduction of the double bond with a suitable reducing agent such as diborane, and oxidation of the primary alcohol to a carboxylic acid with a suitable oxidising agent such as chromium (VI) oxide.

35 Compounds of formula (III) wherein A represents a group (CH₂)₁NR⁹ may be prepared by the reduction of a compound of formula (XXII):



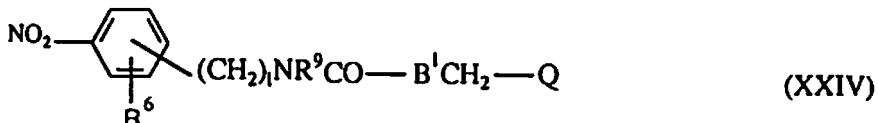
45 (in which B¹ is a bond or a C₁-3 alkylene chain optionally substituted by a hydroxyl group) with a suitable reducing agent such as lithium aluminium hydride in an inert solvent such as an ether (e.g. tetrahydrofuran) at an elevated temperature.

Compounds of formula (XXII) may be prepared by the reduction of a compound of formula (XXIII):



by catalytic hydrogenation, for example as described above for preparing compounds of formula (VI). Compounds of formula (XXIII) may be prepared by the reaction of a compound of formula (XXIV):

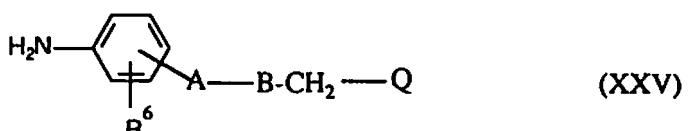
5



10 [wherein Q represents a halogen (e.g. chlorine) atom] with a compound of formula (V) as defined previously under the conditions described above in process (B).

Compounds of formula (IV) may be prepared by the reaction of a compound of formula (II) as defined previously, with a compound of formula (XXV):

15



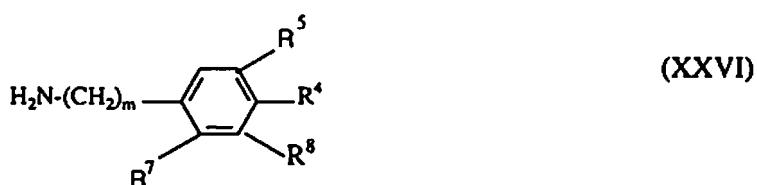
20

wherein Q represents a halogen (e.g. bromine) atom, under the conditions described in process (A) above for the reaction of a compound of formula (II) with a compound of formula (III).

Compounds of formula (V) wherein R³ represents a C₁₋₄alkyl group may be prepared by reacting a compound of formula (XXVI):

25

30



35 with benzaldehyde, followed by a C₁₋₄alkyl halide. Hydrolysis of the resultant quaternary salt followed by treatment with dilute sodium hydroxide solution gives a compound of formula (V) wherein R³ represents a C₁₋₄alkyl group.

It is to be understood that the general procedures above may be used to provide a compound of formula (I) in which B contains a hydroxyl substituent. However, it may be preferable to reduce an intermediate in which B contains an oxo group to provide the desired intermediate in which B contains a hydroxyl substituent at an appropriate stage in the overall procedure.

Intermediates of formulae (III), (IV), (VI), (VII), (VIII), (IX), (XII), (XIII), (XV), (XVI), (XVIII), (XIX), (XXII) and (XXIII) are novel compounds and represent a further aspect of the present invention.

Compounds of formula (II) are either known, or may be prepared by conventional methods, such as those described by G.W.Rewcastle and W.A.Denny in *Synth. Commun.*, 1985, 217-222.

45 Compounds of formulae (V), (IX), (XI), (XII), (XIV), (XVII), (XX), (XXI), (XXIV) and (XXVI) are either known, or may be prepared by conventional methods.

Compounds of formula (XXV) are either known or may be prepared by conventional methods. Thus, for example, compounds of formula (XXV) wherein A represents an oxygen atom may be prepared by the reaction of a 4-acetamidophenol derivative with a dihaloalkane Q-BCH₂-Q, followed by acid hydrolysis using, for example, dilute hydrochloric acid.

Where it is desired to isolate a compound of the invention as a salt, for example a physiologically acceptable salt, this may be achieved by reacting the compound of formula (I) in the form of the free base with an appropriate acid, preferably with an equivalent amount, in a suitable solvent such as an alcohol (e.g. ethanol or methanol), an aqueous alcohol (e.g. aqueous ethanol), a halogenated hydrocarbon (e.g. dichloromethane), an ester (e.g. ethyl acetate) or an ether (e.g. tetrahydrofuran), or a mixture of two or more of such solvents.

Physiologically acceptable salts may also be prepared from other salts, including other physiologically acceptable salts, of the compound of formula (I) using conventional methods.

It will be appreciated that within the above multi-stage processes, the various methods described for the introduction of the desired groups required in the final product may be performed in sequences different from those described. The sequence of the reactions in multi-stage processes should of course be chosen so that the reaction conditions used do not affect groups in the molecule which are desired in the final product.

The invention is further illustrated by the following Intermediates and Examples which are not intended to limit the invention in any way. All temperatures are in °C. ¹H NMR spectra were obtained for dilute solutions in CDCl₃ unless otherwise stated. Solvents were dried, where indicated, over sodium sulphate. Silica gel used for column chromatography was Merck 60, 230-400 mesh. The following abbreviations are used: THF - tetrahydrofuran; DMF - dimethylformamide.

Intermediate 1

(a) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(4-nitrophenoxy)propyl] isoquinoline

A mixture of 1-(3-bromopropoxy)-4-nitrobenzene (10g), 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (8.8g) and potassium carbonate (10.6g) in DMF (100ml) was heated at 100° for 16h. The mixture was then filtered and the filtrate evaporated. The residue was taken up in water and extracted with dichloromethane. The organic layer was washed with water, dried, and evaporated to give an oil which crystallised in ether to give the title compound (11.3g), m.p. 100°.

The following compounds were prepared in a similar manner to Intermediate 1(a):

(b) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-[(4-nitrophenyl)thio]propyl]isoquinoline

The title compound (5.3g) was obtained as an oil (which subsequently crystallised) from 1-[(3-bromopropyl)thio]-4-nitrobenzene (7.0g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (5.8g).
NMR includes d 4.05(6H,s, 2 x OCH₃).

(c) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[2-(4-nitrophenyl)ethyl]isoquinoline

The title compound (16g) was obtained as a solid from 1-(2-bromoethyl)-4-nitrobenzene (10g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (10.9g). M.p. 118°.
NMR includes d 3.9 (6H,s, 2 x OCH₃).

(d) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[4-(4-nitrophenyl)butyl]-isoquinoline

The title compound (12.6g) was obtained as an oil from 1-(4-bromobutyl)-4-nitrobenzene (12.5g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (11.1g). The product was purified by column chromatography eluting with dichloromethane:methanol (99:1).
NMR includes d 3.85 (6H,s, 2 x OCH₃).

Intermediate 2

(a) 4-[3-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy] benzenamine

A solution of Intermediate 1(a) (16g) in ethanol (200ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium on carbon (1.6g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (14.7g) as an oil which crystallised in hexane, m.p. 100°.

(b) 4-[[3-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny) propyl]thio]benzenamine

Intermediate 1(b) (5.3g) was dissolved in a mixture of methanol and concentrated hydrochloric acid (5ml) at room temperature with stirring. Iron powder (3.8g) was then added portionwise, and the mixture was heated under reflux for 1.5h. The mixture was then cooled, poured onto ice, basified with sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water, dried and evaporated to give the title compound (4.35g) as an oil.

IR: Freq NH₂: 3350cm⁻¹.

(c) 4-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]-benzenamine

5 Intermediate 1(c) (14g) was reduced according to the method of Intermediate 2(b) to give the title compound (12g) as a solid, m.p. 120°.

(d) 4-[4-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)butyl]-benzenamine

10 Intermediate 1(d) (8.5g) was reduced according to the method of Intermediate 2(a). The product was purified by column chromatography eluting with dichloromethane: methanol (99:1) to give the title compound (4.3g) as an oil which solidified.

IR: Freq NH₂: 3350 cm⁻¹.

15 Intermediate 3

(a) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[(4-nitrophenoxy)acetyl] isoquinoline

A mixture of (4-nitrophenoxy)acetic acid (50g) and thionyl chloride (150ml) was heated under reflux for 20 3h. The solution was concentrated and then coevaporated with benzene to give 4-nitrophenoxyacetyl chloride as a solid. A solution of this solid (9.4g) in acetone (100ml) was added dropwise to a stirred mixture of 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (10g) and sodium hydrogen carbonate (9g) in acetone (100ml) at 0°. Stirring was continued at room temperature for 16h, the mixture was then filtered, and the filtrate was concentrated. The residue was treated with water and extracted with dichloromethane.

25 The organic layer was washed with water, dried and concentrated to give the title compound (6.6g) as an oil.

IR: Freq CO: 1650cm⁻¹.

The following compound was prepared in a similar manner to Intermediate 3(a).

30 (b) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(4-nitrophenyl)-1-oxopropyl]isoquinoline

The title compound (12.3g) was obtained as a solid, m.p. 134° from 4-nitrobenzenepropanoic acid (9.75g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (11.6g).

35 Intermediate 4

(a) 2-[(4-Aminophenoxy)acetyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

Intermediate 3(a) (6.6g) was dissolved in a mixture of methanol (100ml) and concentrated hydrochloric acid (50ml) at room temperature with stirring. Iron powder (5g) was then added portionwise and the mixture was heated under reflux for 3h. The mixture was then cooled, poured onto ice, basified with sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water, dried and evaporated to give the title compound (4g) as an oil.

IR: Freq NH₂: 3360cm⁻¹.

45 (b) 2-[3-(4-Aminophenyl)-1-oxopropyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

A solution of Intermediate 3(b) (12g) in a mixture of ethanol:dioxan (18ml; 5:1) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium on carbon (1.2g). After 50 hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (11g) as a solid.

IR: Freq NH₂: 3360cm⁻¹

Freq CO: 1650cm⁻¹.

55 Intermediate 5

(a) 4-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethoxy] benzenamine

A solution of Intermediate 4(a) (4g) in THF (50ml) was added dropwise to a stirred suspension of lithium aluminium hydride (1.8g) in THF (20ml) at room temperature, and the mixture was heated under reflux for 3h. Water was added carefully to the cooled mixture which was then filtered, washed with THF, evaporated and extracted with dichloromethane. The organic layer was dried and evaporated to give the title compound (1.5g) as an oil.

IR: Freq NH₂: 3350cm⁻¹.

The following compound was prepared in a similar manner to Intermediate 5(a):

(b) 4-[3-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl] benzenamine

The title compound (8.6g) was obtained as a solid, m.p. 138°, by the reduction of Intermediate 4(b) (11g).

Intermediate 6

(a) 1-(3-Bromopropoxy)-3-methoxy-4-nitrobenzene

A mixture of Intermediate 18 (2.4g), 1,3-dibromopropane (7.5ml) and potassium carbonate (2.2g) in DMF (30ml) was stirred at room temperature for 24h. The mixture was filtered and the filtrate was evaporated to dryness. The residue was treated with water and extracted with dichloromethane. The organic extract was then washed with 5% sodium hydroxide solution and brine, dried and concentrated in vacuo to give the title compound (3.5g) as an oil.

NMR includes d 2.3 (2H,m,CH₂), 3.6 (2H,t,CH₂Br), 3.8 (3H,s,OCH₃), 4.1 (2H,t, CH₂O).

The following compounds were prepared in a similar manner to Intermediate 6(a):

(b) 1-(3-Bromopropoxy)-3-methyl-4-nitrobenzene

The title compound (33g) was obtained as an oil from 3-methyl-4-nitrophenol (25g) and 1,3-dibromopropane (83ml).

NMR includes d 2.3 (2H,m,CH₂), 2.5 (3H,s,CH₃), 3.6 (2H,t,CH₂Br), 4.1 (2H,t,OCH₂).

(c) 1-(3-Bromopropoxy)-3-ethyl-4-nitrobenzene

The title compound was obtained from 3-ethyl-4-nitrophenol and 1,3-dibromopropane. NMR includes d 1.23 (t,3H,CH₃-CH₂-), 2.2 (m,2H,CH₂-CH₂-CH₂), 2.8 (q,2H,CH₂-CH₃), 3.5 (t,2H,CH₂Br), 4.1 (t,2H,O-CH₂-), 6.6 (m,2H,Ar), 7.8 (d,2H,Ar).

Intermediate 7

(a) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(3-methoxy-4-nitrophenoxy)propyl]isoquinoline

A mixture of Intermediate 6(a) (0.7g), 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (0.4g) and potassium carbonate (0.36g) in DMF (25ml) was heated at 60° for 16h. The mixture was filtered and the filtrate was evaporated. The residue was treated with water and extracted with dichloromethane. The organic layer was dried, concentrated, and the resultant residue was purified by column chromatography eluting with dichloromethane:methanol (99:1) to give the title compound (0.64g) as an oil.

NMR includes d 3.8 (9H,2s, 3 x OCH₃).

The following compound was prepared in a similar manner to Intermediate 7(a):

(b) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(3-methyl-4-nitrophenoxy)propyl]isoquinoline

The title compound (5.3g) was obtained as an oil from Intermediate 6(b) (5.7g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (4.0g).

NMR includes d 2.5 (3H,s,CH₃), 3.8 (6H,s, 2 x OCH₃)

Intermediate 8

(a) 2-Methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]benzenamine

A solution of Intermediate 7(a) (0.64g) in ethanol (25ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium on carbon (60mg). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated in vacuo to give the title compound (0.4g) as a solid.

5 NMR includes d 3.8 (9H,s, 3 x OCH₃), 3.0 (2H,bs,NH₂).

The following compound was prepared in a similar manner to Intermediate 8(a):

(b) 2-Methyl-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]benzenamine

10 The title compound (4.8g) was obtained as an oil (which subsequently crystallised) from Intermediate 7-(b) (5.3g).

NMR includes d 2.1 (3H,s,CH₃), 3.8 (6H, s, 2 x OCH₃).

Intermediate 9

15

(a) 3-Methyl-4-nitrobenzenoic acid

20 3-Methyl-4-nitrobenzoyl chloride (10g) in ether (100ml) was added dropwise to a solution of diazomethane (prepared from 30g of N-methyl-N-nitroso-p-toluene sulphonamide) at 0°. The reaction mixture was stirred at room temperature for 3h and then concentrated in vacuo to give the diazo ketone as a solid. This diazo ketone in dioxan (100ml) was then added dropwise to a solution of silver oxide in water (prepared from silver nitrate (20g) and dilute sodium hydroxide (100ml)). The mixture was stirred at 75-80° for 3.5h and filtered. The filtrate was diluted with water, acidified with a solution of nitric acid and the product was extracted with hot diisopropyl ether, treated with brine and concentrated in vacuo to give the title compound (6g) as a solid, m.p. 95°.

25

In the same way, the following compound was prepared:

(b) 3-Methoxy-4-nitrobenzenoic acid, m.p. 130-131°.

30

From 3-methoxy-4-nitrobenzoyl chloride.

Intermediate 10

Ethyl 3-(3-hydroxy-4-nitrophenyl)-2-propenoate

35

To a solution of 3-hydroxy-4-nitrobenzaldehyde (5g) in toluene (50ml) was added carbethoxymethylenetriphenylphosphorane (8.96g), and the mixture was heated under reflux for 2h. The mixture was then concentrated and the residue was purified by column chromatography eluting with cyclohexane:ethyl acetate (6:4) to give the title compound (6.2g) as a solid, m.p. 95°.

40

Intermediate 11

Ethyl 3-(3-methoxy-4-nitrophenyl)-2-propenoate

45

To a solution of Intermediate 10 (5.88g) in DMF (50ml) was added potassium carbonate (4.4g) and methyl iodide (4ml). The mixture was stirred at room temperature for 2h and then concentrated in vacuo. The residue was treated with water and extracted with dichloromethane. The organic extract was dried and concentrated to give the title compound (6.2g) as a solid, m.p. 130°.

50

Intermediate 12

3-(3-Methoxy-4-nitrophenyl)-2-propenoic acid

55 To a suspension of Intermediate 11 (6.2g) in ethanol (50ml) was added a solution of 1N sodium hydroxide (50ml). The mixture was heated under reflux for 1h and then poured onto cracked ice. A solution of 1N hydrochloric acid (60ml) was added and the precipitate was filtered off to give the title compound (4g) as a solid. NMR (DMSO-d₆) includes d 3.95 (3H,s,0CH₃).

Intermediate 133-(3-Ethoxy-4-nitrophenyl)-2-propenoic acid

5 Using reactions similar to those described in Intermediates 11 and 12, the title compound (3.1g) was obtained as a solid, m.p. 272°, from Intermediate 10 (4.0g), ethyl iodide (4ml) and potassium carbonate (2.6g), followed by saponification of the ester function.

Intermediate 14

10 (a) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(3-methoxy-4-nitrophenyl)-1-oxo-2-propenyl]isoquinoline

A mixture of Intermediate 12 (4.9g) and 1-hydroxybenzotriazole (2.95g) in DMF (100ml) was stirred at room temperature for 10 min. 1,2,3,4-Tetrahydro-6,7 dimethoxy-isoquinoline (5g) was added, followed by dicyclohexylcarbodiimide (4.52g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute hydrochloric acid, then dilute sodium hydroxide solution and extracted with dichloromethane. The organic extract was dried, concentrated in vacuo, and the residue was purified by column chromatography eluting firstly with ethyl acetate:cyclohexane (4:6), then with ethyl acetate to give the title compound which was crystallised from ethyl acetate/ether and obtained as crystals (6.5g).

NMR includes d 3.85 (6H,s, 2 x OCH₃), 3.95 (3H,s,OCH₃).

The following compounds were prepared in a similar manner to Intermediate 14(a):

(b) 2-[3-(3-Ethoxy-4-nitrophenyl)-1-oxo-2-propenyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

25 The title compound (5.3g) was obtained as a solid, m.p. 152° from Intermediate 13 (3.0g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (2.5g).

(c) 1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[(3-methyl-4-nitrophenyl)acetyl]isoquinoline

30 The title compound (2.8g) was obtained as an oil from Intermediate 9(a) (1.8g) and 1,2,3,4-tetrahydro-6,7-dimethoxy-isoquinoline (1.9g).

IR: Freq CO: 1650cm⁻¹.

Intermediate 15

(a) 2-[3-(4-Amino-3-methoxyphenyl)-1-oxopropyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

A solution of Intermediate 14(a) (6.5g) in methanol/ethyl acetate (1:1; 100ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium on carbon (0.3g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated in vacuo to give the title compound (6g) as an oil.

NMR includes d 3.8 (9H,s, 3 x OCH₃).

The following compounds were prepared in a similar manner to Intermediate 15(a):

(b) 2-[3-(4-Amino-3-ethoxyphenyl)-1-oxopropyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

The title compound (4.5g) was obtained as an oil from Intermediate 14(b) (5.3g).

IR: Freq CO: 1640cm⁻¹

50 Freq NH₂: 3450cm⁻¹.

(c) 2-[(4-Amino-3-methylphenyl)acetyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

The title compound (2.4g) was obtained as an oil from Intermediate 14(c) (2.8g).

55 IR: Freq CO: 1650cm⁻¹

Freq NH₂: 3340-3440cm⁻¹.

Intermediate 16

(a) 2-Methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]benzenamine

5 A solution of Intermediate 15(a) (6g) in THF (30ml) was added dropwise to a stirred suspension of lithium aluminium hydride (1.84g) in THF (50ml) at room temperature, and the mixture was heated under reflux for 2h. Water was carefully added to the cooled mixture which was then filtered. The filtrate was concentrated in vacuo, treated with water and extracted with dichloromethane. The organic layer was dried and concentrated in vacuo to give the title compound (4.2g) as an oil.

IR: Freq NH₂: 3340-3440cm⁻¹.

10 The following compounds were prepared in a similar manner to Intermediate 16(a):

(b) 2-Ethoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]benzenamine

The title compound (2.5g) was obtained as an oil from Intermediate 15(b) (4.5g).

IR: Freq NH₂: 3340-3440cm⁻¹.

(c) 2-Methyl-4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]benzenamine

The title compound (1.7g) was obtained as a solid, m.p. 105°, from Intermediate 15(c) (2.4g).

20 Intermediate 17**3-Chloro 4-nitrophenol**

25 Concentrated nitric acid (10ml) in acetic acid (30ml) was added dropwise to a cooled solution of 3-chlorophenol (10g) in acetic acid (10ml). After 1 hour at - 5°, the mixture was poured onto ice, extracted with ether, dried over sodium sulfate and evaporated. The residue was then purified by column chromatography eluting with hexane-ethyl acetate (85:15) to give the title compound (9g). M.p. 120°.

Intermediate 18**3-Methoxy-4-nitrophenol**

35 A solution of Intermediate 17 (4.4g) in methanol (15ml) was added to a solution of sodium (5.8g) in methanol (60ml) and the mixture was stirred in an autoclave for 16 h at 100°. The mixture was cooled and poured onto ice and acidified with concentrated hydrochloric acid. Methanol was then evaporated in vacuo inducing the crystallisation of the title compound (3.5g). M.p. 142°.

Intermediate 19**40 1-(2-Chloroethoxy)-3-methyl-4-nitrobenzene**

45 A mixture of 3-methyl-4-nitrophenol (10g), 1-bromo-2-chloroethane (16ml) and sodium hydroxide (2.9g) in water (50 ml) was stirred under reflux for 16h. The mixture was diluted with water and the product was extracted with methylene chloride. The organic extract was dried on sodium sulfate and concentrated in vacuo to give the title compound as an oil (10.81g). NMR includes d 2.5 (s,3H,-CH₃), 3.9 (t,2H,CH₂-O) and 4.3 (t,2H,-CH₂-Cl).

Intermediate 20**50 (a) 3,4-Dimethoxy-N-methylbenzenethanamine**

3,4-Dimethoxybenzenethanamine (100g) was mixed with benzaldehyde (59g), and rotoevaporated to give an oil. Methyl iodide (69 ml) was then added and the mixture was heated for 48h at 40° and then boiled with 80% ethanol (500ml) for 3h. After half of the ethanol had evaporated, the solution was treated with ether (1 litre) to give a solid that was filtered, washed with ether, treated with dilute sodium hydroxide and extracted with ether to give the title compound (80g) as an oil that was distilled under reduced pressure, b.p. 0.1mm; 92-95°.

(b) 3,4-Dimethoxy-N-methylbenzenemethanamine

3,4-Dimethoxybenzenemethanamine (100 g) was mixed with benzaldehyde (64g), and rotoevaporated to give an oil. Methyl iodide (75 ml) was then added and the mixture was heated for 48h at 40° and then 5 boiled with 80% ethanol (800ml) for 3h. After half of the ethanol had evaporated, the solution was treated with ether (1 litre) to give a solid that was filtered, washed with ether, treated with dilute sodium hydroxide and extracted with ether to give the title compound (69g) as an oil that was distilled under reduced pressure, b.p. 0.03mm; 91°.

The following amines were prepared in a similar manner to Intermediates 20(a) and 20(b):

10

(c) 4-Fluoro-N-methylbenzenemethanamine as an oil; IR includes a peak at 3300cm⁻¹ (NH).

From 4-fluorobzenemethanamine and methyl iodide.

15

(d) 4-Methoxy-N-methylbenzenemethanamine as an oil; IR includes a peak at 3310cm⁻¹ (NH).

From 4-methoxybenzenemethanamine and methyl iodide.

20

(e) 4-Methoxy-N-methylbenzeneethanamine as an oil; IR includes a peak at 3310cm⁻¹ (NH).

From 4-methoxybenzeneethanamine and methyl iodide.

25

(f) 4-(Methylthio)-N-methylbenzenemethanamine as an oil; IR includes a peak at 3310cm⁻¹ (NH).

From 4-(methylthio)benzenemethanamine and methyl iodide.

30

(g) 4-Methyl-N-Methylbenzenemethanamine as an oil; IR includes a peak at 3310cm⁻¹ (NH).

From 4-methylbenzenemethanamine and methyl iodide.

35

Intermediate 21(a) 3,4-Dimethoxy-N-methyl-N-[3-(3-methyl-4-nitrophenoxy)propyl] benzenemethanamine

40

A mixture of Intermediate 6(b) (6g), Intermediate 20(b) (4g) and potassium carbonate (3.3g) in DMF (80ml) was heated at 60° for 36h. The mixture was filtered and the filtrate was evaporated. The residue was added to water and extracted with dichloromethane. The organic layer was washed with water, dried over sodium sulfate filtered and evaporated. The oily residue was then chromatographed with dichloromethane/methanol (99:1) to give the title compound as an oil (4.6 g). NMR includes d 2.2 (s,3H,-CH₃), 2.4 (s,3H,N-CH₃) and 3.8 (s,6H,2OCH₃).

In the same way, the following compounds were prepared:

45

(b) 3,4-Dimethoxy-N-[3-(3-methoxy-4-nitrophenoxy)propyl]-N-methylbenzenemethanamine as an oil

50

From Intermediate 6(a) and Intermediate 20(b). NMR includes d 2.2 (s,3H,N-CH₃) and 3.85 - 3.9 (2s,3H,6H,3O-CH₃).

55

(c) 3,4-Dimethoxy-N-[3-(3-ethyl-4-nitrophenoxy)propyl]-N-methylbenzenemethanamine as an oil.

60

From Intermediate 6(c) and Intermediate 20(b). NMR includes d 2.2 (s,3H,N-CH₃) and 3.85 - 3.9 (s,6H,2O-CH₃).

65

(d) 3,4-Dimethoxy-N-methyl-N-[2-(3-methyl-4-nitrophenoxy)ethyl] benzenemethanamine as an oil

70

From Intermediate 19 and Intermediate 20(b). NMR includes d 2.3 (s,3H,N-CH₃), 2.5 (s,3H,N-CH₃) and 3.8 (s,6H,2OCH₃).

Intermediate 22

(a) N-[3-(4-Amino-3-methylphenoxy)propyl]-3,4-dimethoxy-N-methylbenzenemethanamine

5 A solution of Intermediate 21(a) (4.6g) in ethanol (100ml) was hydrogenated at room temperature in presence of 10% palladium-on-carbon 10% (450mg). After the hydrogen absorption was completed, the catalyst was filtered off and the solution concentrated to give the title compound (3.7g) as an oil. NMR includes d 2.0 (s,3H,CH₃), 2.1 (s,3H,N-CH₃) and 3.7 (s,6H,2OCH₃).

In the same way, the following compounds were prepared:

(b) N-[3-(4-Amino-3-methoxyphenoxy)propyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil.

10 From Intermediate 21(b). NMR includes d 2.2 (s,3H,N-CH₃), 3.85-3.9 (s,3H,OCH₃) and 3.9 (s,6H,2OCH₃).

(c) N-[3-(4-Amino-3-ethylphenoxy)propyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil.

15 From Intermediate 21(c). NMR includes d 2.1 (s,3H,N-CH₃) and 3.7 (s,6H,2OCH₃).

(d) N-[2-(4-Amino-3-methylphenoxy)ethyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil.

20 From Intermediate 21(d). NMR includes d 2.0 (s,3H,N-CH₃), 2.2 (s,3H,N-CH₃) and 3.8 (s,6H,2OCH₃).

Intermediate 23Diethyl (3-methyl-4-nitrobenzyl)malonate

25 To a solution of sodium ethanolate [prepared from 1.35g Na in ethanol (30ml)] were added diethyl malonate (9.2ml) and then dropwise 3-methyl-4-nitrobenzyl bromide (13.4g). The mixture was stirred 30 minutes at room temperature, then 30 minutes under reflux and then concentrated. The residue is treated with water and hexane, the precipitate filtered and the filtrate extracted with diethyl ether. The organic extract was dried on sodium sulfate and concentrate to give the title compound as an oil (4g).

30 NMR includes d 1.15 (t,6H,2xCH₃-CH₂), 2.5 (s,3H,CH₃-Ar), 3.16 (s,2H,CH₂-Ar), 4.0 (q,4H,2xCH₂-CH₃), 7.0 (m,2H,Ar), 7.7 (d,1H,Ar).

Intermediate 243-(3-Methyl-4-nitrophenyl)propionic acid

35 Intermediate 23 (4g) was added dropwise to a solution of potassium hydroxide (3.1g) in water and the mixture is stirred under reflux for 2 hours, diluted with water, washed with diethyl ether and then acidified with a dilute solution of hydrochloric acid. After extraction with diethyl ether and concentration, the 40 concentrate was heated at 130° for 3h to give the title compound as a yellow solid (2.3g). NMR (CDCl₃) includes d 2.5 (s,3H,CH₃) and 2.9 (m,4H,2CH₂).

Intermediate 25(a) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-methyl-4-nitrobenzenethanamide

45 A mixture of Intermediate 9(a) (2g) and 1-hydroxybenzotriazole (1.6g) in DMF (35ml) was stirred at room temperature for 5 min. Intermediate 20(b) (1.9g) in DMF (20 ml) was then added, followed by dicyclohexyl-carbodiimide (2.1g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate 50 was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated and the residue was purified by column chromatography eluting with dichloromethane/methanol (97:3) to give the title compound (1.7g) as an oil. IR includes a signal at 1640cm⁻¹(CO).

In the same way, the following compounds were prepared:

(b) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-methoxy-4-nitrobenzenethanamide

55 From Intermediate 9(b) and Intermediate 20(b). IR includes a signal at 1645cm⁻¹ (CO).

(c) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-methyl-4-nitrobenzenepropanamide as an oil

From Intermediate 24 and Intermediate 20(b). NMR (CDCl_3) includes d 2.5 (s,3H,-CH₃), 2.9 (s,3H,N-CH₃) and 3.8 (s,6H,2OCH₃).

5

Intermediate 26(a) 4-Amino-3-methyl-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzeneethanamide

10 A solution of Intermediate 25(a) (1.7g) in ethanol (60ml) was hydrogenated at room temperature in presence of 10% palladium-on- carbon (0.25g). After the hydrogen absorption was completed the catalyst was filtered off and the solution concentrated to give the title compound (1.4g) as an oil. IR includes signals at 3450-3350 cm⁻¹ (NH₂) and 1630 cm⁻¹ (CO).

In the same way, the following compounds were prepared:

15

(b) 4-Amino-3-methoxy-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzeneethanamide

From Intermediate 25(b). IR includes signals at 3450-3350cm⁻¹ (NH₂) and 1625 cm⁻¹ (CO).

20

(c) 4-Amino-3-methyl-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamide

From Intermediate 25(c). NMR includes d 2.1 (3H,s,CH₃), 2.75 (3H,s,N-CH₃) and 3.8 (6H,s,2OCH₃).

Intermediate 27

25

(a) 4-Amino-3-methyl-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzeneethanamine

A solution of Intermediate 26(a) (1.4g) in THF (50ml) was added dropwise to a stirred suspension of lithium aluminium hydride (0.7g) in THF (30 ml) at room temperature and the mixture was heated under reflux for 3h. Water was added carefully to the cooled mixture which was then filtered on a celite pad, washed with THF, evaporated and extracted with ether. The ethereal extracts were dried and evaporated to give the title compound (1g) as an oil. IR includes a signal at 3450 - 3350 cm⁻¹ (NH₂).

In the same way, the following compounds were prepared:

35

(b) 4-Amino-3-methoxy-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzeneethanamine

From Intermediate 26(b). IR includes a signal at 3455 - 3345 cm⁻¹ (NH₂).

40

(c) 4-Amino-3-methyl-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamine as an oil

From Intermediate 26(c). NMR includes d 2.0 (3H,s,-CH₃), 2.1 (3H,s,N-CH₃) and 3.8 (6H,s,2OCH₃).

Intermediate 28

45

N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-methoxy-4-nitrobenzene-2-propenamide

A mixture of Intermediate 12 (3g) and 1-hydroxybenzotriazole (1.95g) in DMF (100 ml) was stirred at room temperature for 10 minutes. Intermediate 20(b) (2.5g) was added, followed by dicyclohexylcarbodiimide (2.95g) and the mixture was stirred at room temperature for 16 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute hydrochloric acid solution, then dilute sodium hydroxide solution and extracted with methylene chloride. The organic extract was dried with sodium sulfate and concentrated. The residue was purified by column chromatography eluting with ethyl acetate to give the title compound (4.4g).

NMR includes d 2.9 (3H,s,N-CH₃), 3.85 (3H,s,OCH₃) and 3.9 (6H,s,2OCH₃).

55

Intermediate 294-Amino-3-methoxy-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamide

A solution of Intermediate 28 (8.4g) in methanol/ethyl acetate (1:1, 100ml) was hydrogenated at room temperature in presence of 10% palladium-on-carbon (0.3g). After the hydrogen absorption was completed, the catalyst was filtered off and the solution concentrated to give the title compound (7.3g) as an oil. IR includes signals at 3450-3350 cm⁻¹ (NH₂) and 1635 cm⁻¹ (CO).

5

Intermediate 30

4-Amino-3-methoxy-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamine

10 A solution of Intermediate 29 (7.32g) in tetrahydrofuran (100 ml) was added dropwise to a stirred suspension of lithium aluminium hydride (2.3g) in tetrahydrofuran (100 ml) at room temperature and the mixture was heated under reflux 1 h. Water (20 ml) was added carefully to the cooled mixture which was filtered on a celite pad, washed with diethyl ether, concentrated and extracted with methylene chloride. The organic extract was dried on sodium sulfate, evaporated and the product purified by column chromatography on silica gel eluting with dichloromethane/methanol (95:5) to give the title compound as an oil (2.5g). IR includes a signal at 3440-3340 cm⁻¹ (NH₂).

Intermediate 31

20 (a) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-nitrobenzenebutanamide

A mixture of 4-nitrobenzenebutanoic acid (31 g) and thionyl chloride (200ml) was heated under reflux for 1 h. The solution was then concentrated and coevaporated with benzene to give an oil. This oil was dissolved in acetone (100ml) and added dropwise to a stirred mixture of Intermediate 20(b) (28.6g) and 25 sodium hydrogen carbonate (35 g) in acetone (150 ml) at room temperature. Stirring was continued for 4h, the mixture was then filtered and the filtrate was concentrated. The residue was poured into water and then extracted with dichloromethane. The organic phase was evaporated to give the title compound (41.5 g) as an oil. Recrystallisation from ethanol gave the title compound as a solid, MP : 90°.

30 (b) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-nitrobenzeneethanamide

A mixture of 4-nitrobenzeneacetic acid (22 g) and thionyl chloride (200 ml) was heated under reflux for 3h. The solution was concentrated and then coevaporated with benzene to give an oil. This oil was dissolved in acetone (100ml) and added dropwise to a stirred mixture of Intermediate 20(b) (22g) and 35 sodium hydrogen carbonate (15.3 g) in acetone (100 ml) at room temperature. Stirring was continued for 6 hours, the mixture was then filtered and the filtrate was concentrated. The residue was poured into water and extracted with ethyl acetate. The organic phase was washed first with dilute sodium hydroxide solution, then with water, dried and concentrated to give the title compound (22.3g) as an oil. IR includes a peak at 1650cm⁻¹ (CO).

40 The following amides were prepared in a similar manner to Intermediates 31(a) and 31(b):

(c) N-[2-(3,4-Dimethoxyphenyl)ethyl]-N-methyl-4-nitrobenzenebutanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

45 From 4-nitrobenzenebutanoic acid and Intermediate 20(a).

(d) N-[2-(3,4-Dimethoxyphenyl)ethyl]-N-methyl-4-nitrobenzenepropanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

50 From 4-nitrobenzenepropanoic acid and Intermediate 20(a).

(e) N-[2-(3,4-Dimethoxyphenyl)ethyl]-N-methyl-4-nitrobenzeneethanamide as an oil; IR includes a peak at 1650cm⁻¹ (CO).

55 From 4-nitrobenzeneacetic acid and Intermediate 20(a).

(f) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-nitrobenzenepropanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

From 4-nitrobenzenepropanoic acid and Intermediate 20(b).

(g) N-[(4-Methoxyphenyl)methyl]-N-methyl-4-nitrobenzenepropanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

5

From 4-nitrobenzenepropanoic acid and Intermediate 20(d).

(h) N-[2-(4-Methoxyphenyl)ethyl]-N-methyl-4-nitrobenzenebutanamide as an oil; IR includes a peak at 1650cm⁻¹ (CO).

10

From 4-nitrobenzenebutanoic acid and Intermediate 20(e).

(i) N-[(4-Fluorophenyl)methyl]-N-methyl-4-nitrobenzenebutanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

15

From 4-nitrobenzenebutanoic acid and Intermediate 20(c).

(j) N-[(4-(Methylthio)phenyl)methyl]-N-methyl-4-nitrobenzenebutanamide as an oil; IR includes a peak at 1640cm⁻¹ (CO).

20

From 4-nitrobenzenebutanoic acid and Intermediate 20(f).

(k) N-[2-(4-Methoxyphenyl)ethyl]-N-methyl-4-nitrobenzenethanamide as an oil; IR includes a peak at 1650cm⁻¹ (CO).

25

From 4-nitrobenzeneacetic acid and Intermediate 20(e).

(l) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-nitrobenzenepentanamide as an oil; IR includes a peak at 1650cm⁻¹ (CO).

30

From 4-nitrobenzenepentanoic acid and Intermediate 20(b).

Intermediate 32

35 (a) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenebutanamide

Intermediate 31(a) (40g) was dissolved in a mixture of methanol (300 ml) and concentrated hydrochloric acid (160ml) at room temperature with stirring. Iron powder (21 g) was then added slowly, and the reaction mixture was heated under reflux for 1h. The mixture was then evaporated and basified with sodium hydroxide solution. Ethyl acetate (1 litre) was added and the mixture was filtered. The organic phase was washed with water, dried and evaporated to give the title compound (30 g) as an oil. IR includes peaks at 1630 cm⁻¹ (CO), 3350-3430cm⁻¹ (NH₂).

40 (b) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenethanamide

45

Intermediate 31(b) (22g) was dissolved in a mixture of methanol (300 ml) and concentrated hydrochloric acid (150 ml) at room temperature with stirring. Iron powder (18 g) was then added slowly, and the reaction mixture was heated under reflux for 3 h. The mixture was then evaporated, basified with sodium hydroxide solution, and extracted with ethyl acetate. The organic phase was washed with water, dried and evaporated to give the title compound (14 g) as an oil. IR includes peaks at 1620cm⁻¹ (CO) and 3350-3450cm⁻¹ (NH₂).

The following compounds were prepared in a similar manner to Intermediates 32(a) and 32(b):

(c) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzenebutanamide as an oil; IR includes peaks at 1630cm⁻¹ (CO) and 3330-3420cm⁻¹ (NH₂).

55

From Intermediate 31(c).

(d) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzenepropanamide as an oil; IR includes peaks at

1630cm⁻¹ (CO) and 3340-3420cm⁻¹ (NH₂).

From Intermediate 31(d).

5 (e) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzeneethanamide as an oil; IR includes peaks at 1640cm⁻¹ (CO) and 3330-3420cm⁻¹ (NH₂).

From Intermediate 31(e).

10 (f) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamide as an oil; IR includes peaks at 1640cm⁻¹ (CO) and 3350-3440cm⁻¹ (NH₂).

From Intermediate 31(f).

15 (g) 4-Amino-N-[(4-methoxyphenyl)methyl]-N-methylbenzenepropanamide as an oil; IR includes peaks at 1650cm⁻¹ (CO) and 3330-3420cm⁻¹ (NH₂).

From Intermediate 31(g).

20 (h) 4-Amino-N-[2-(4-methoxyphenyl)ethyl]-N-methylbenzenebutanamide as an oil; IR includes peaks at 1640cm⁻¹ (CO) and 3340-3430cm⁻¹ (NH₂).

From Intermediate 31(h).

25 (i) 4-Amino-N-[(4-fluorophenyl)methyl]-N-methylbenzenebutanamide as an oil; IR includes peaks at 1640cm⁻¹ (CO) and 3340-3430cm⁻¹ (NH₂).

From Intermediate 31(i).

30 (j) 4-Amino-N-[(4-(methylthio)phenyl)methyl]-N-methylbenzenebutanamide as an oil; IR includes peaks at 1640cm⁻¹ (CO) and 3340-3430cm⁻¹ (NH₂).

From Intermediate 31(j).

35 (k) 4-Amino-N-[2-(4-methoxyphenyl)ethyl]-N-methylbenzeneethanamide as an oil; IR includes peaks at 1635cm⁻¹ (CO) and 3340-3440cm⁻¹ (NH₂).

From Intermediate 31(k).

40 (l) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepentanamide as an oil; IR includes peaks at 1630cm⁻¹ (CO) and 3340-3420cm⁻¹ (NH₂).

From Intermediate 31(l).

45 Intermediate 33

(a) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenebutanamine

A solution of Intermediate 32(a) (30g) in THF (150 ml) was added dropwise to a stirred suspension of lithium aluminium hydride (10g) in THF (150 ml) at room temperature and the mixture was heated under reflux for 3h. Water was added carefully to the cooled mixture, which was then filtered, washed with THF, evaporated, and extracted with ether. The combined ethereal extracts were dried and evaporated to give the title compound (21 g) as an oil. IR includes a peak at 3370-3440cm⁻¹ (NH₂).

55 (b) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzeneethanamine

A solution of Intermediate 32(b) (14g) in THF (100 ml) was added dropwise to a stirred suspension of lithium aluminium hydride (8 g) in THF (100 ml) at room temperature and the mixture was heated under

reflux for 3 hours. Water was added carefully to the cooled mixture which was then filtered, washed with THF, evaporated and extracted with ether. The combined ethereal extracts were dried and evaporated to give the title compound (9.5 g) as an oil. IR includes a peak at 3360-3430cm⁻¹ (NH₂).

The following compounds were prepared in a similar manner to Intermediates 33(a) and 33(b):

5 (c) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzenebutanamine as an oil; IR includes a peak at 3360-3430cm⁻¹ (NH₂).

From Intermediate 32(c).

10 (d) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzenepropanamine as an oil; IR includes a peak at 3360-3460cm⁻¹ (NH₂).

From Intermediate 32(d).

15 (e) 4-Amino-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylbenzenethanamine as an oil; IR includes a peak at 3360-3430cm⁻¹ (NH₂).

From Intermediate 32(e).

20 (f) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamine as an oil; IR includes a peak at 3360-3440cm⁻¹ (NH₂).

From Intermediate 32(f).

25 (g) 4-Amino-N-[(4-methoxyphenyl)methyl]-N-methylbenzenepropanamine as an oil; IR includes a peak at 3360-3430cm⁻¹ (NH₂).

From Intermediate 32(g).

30 (h) 4-Amino-N-[2-(4-methoxyphenyl)ethyl]-N-methylbenzenebutanamine as an oil; IR includes a peak at 3380-3460cm⁻¹ (NH₂).

From Intermediate 32(h).

35 (i) 4-Amino-N-[(4-fluorophenyl)methyl]-N-methylbenzenebutanamine as an oil; IR includes a peak at 3350-3430cm⁻¹ (NH₂).

From Intermediate 32(i).

40 (j) 4-Amino-N-[[4-(methylthio)phenyl]methyl]-N-methylbenzenebutanamine as an oil; IR includes a peak at 3350-3430cm⁻¹ (NH₂).

From Intermediate 32(j).

45 (k) 4-Amino-N-[2-(4-methoxyphenyl)ethyl]-N-methylbenzenethanamine as an oil; IR includes a peak at 3360-3440cm⁻¹ (NH₂).

From Intermediate 32(k).

50 (l) 4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepentanamine as an oil; IR includes a peak at 3360-3440cm⁻¹ (NH₂).

From Intermediate 32(l).

55 Intermediate 34

(a) N-[2-(3,4-Dimethoxyphenyl)ethyl]-N-methyl-2-(4-nitrophenoxy)acetamide

A mixture of (4-nitrophenoxy)acetic acid (51 g) and thionyl chloride was heated under reflux for 2h. The solution was concentrated and then coevaporated with benzene to give a solid. This solid was dissolved in acetone (250 ml) and added dropwise to a stirred mixture of Intermediate 20(a) (50g) and sodium hydrogen carbonate (22g) in acetone (250 ml) at room temperature. Stirring was continued for 4h, the mixture was 5 then filtered and the filtrate was concentrated. The residue was treated with water and extracted with ethyl acetate. The organic phase was washed first with dilute sodium hydroxide, then with water, dried and concentrated. Recrystallisation from ethanol gave the title compound (82 g). MP 121°.

The following compounds were prepared in a similar manner to Intermediate 34(a):

10 (b) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-2-(4-nitrophenoxy)acetamide. MP 130°
 From (4-nitrophenoxy)acetic acid and Intermediate 20(b).

(c) N-Methyl-2-(4-nitrophenoxy)-N-(phenylmethyl)acetamide. MP 98°.
 15 From (4-nitrophenoxy)acetic acid and N-methylbenzenemethanamine.

(d) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-2-(4-nitrophenoxy)acetamide as an oil. NMR includes signals at δ 3.0 (3H,s,N-CH₃) and 3.8 (6H,s,OCH₃).
 20 From (4-nitrophenoxy)acetic acid and Intermediate 20(b).

(e) N-[2-(4-Methoxyphenyl)ethyl]-N-methyl-2-(4-nitrophenoxy)acetamide. MP 107°.
 25 From (4-nitrophenoxy)acetic acid and Intermediate 20(e).

(f) N-[(4-Methoxyphenyl)methyl]-N-methyl-2-(4-nitrophenoxy)acetamide. MP 120°.
 From (4-nitrophenoxy)acetic acid and Intermediate 20(d).

30 (g) N-Methyl-N-[(4-methylphenyl)methyl]-2-(4-nitrophenoxy)acetamide. MP 126°.
 From (4-nitrophenoxy)acetic acid and Intermediate 20(g).

35 (h) N-Methyl-N-[(4-(methylthio)phenyl)methyl]-2-(4-nitrophenoxy)acetamide. MP 122°.
 From (4-nitrophenoxy)acetic acid and Intermediate 20(f).
 (i) N-Ethyl-2-(4-nitrophenoxy)-N-(phenylmethyl)acetamide as an oil; IR includes a peak at 1655cm⁻¹ (CO).
 40 From (4-nitrophenoxy)acetic acid and N-ethylbenzenemethanamine.

Intermediate 35

45 (a) 2-(4-Aminophenoxy)-N-[2-(3,4-dimethoxyphenyl)ethyl]-N-methylacetamide
 A solution of Intermediate 34(a) (37.5g) in ethanol (350 ml) was hydrogenated at room temperature in the presence of 10% palladium on carbon (3.5 g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (34 g) as an oil. IR includes 50 peaks at 1650cm⁻¹ (CO) and 3340-3400cm⁻¹ (NH₂).
 The following compounds were prepared in a similar manner to Intermediate 35(a):

(b) 2-(4-Aminophenoxy)-N-[(3,4-dimethoxyphenyl)methyl]-N-methylacetamide as an oil. IR includes peaks at 1650cm⁻¹ (CO) and 3340-3400cm⁻¹ (NH₂).
 55 From Intermediate 34(b).

(c) 2-(4-Aminophenoxy)-N-methyl-N-(phenylmethyl)acetamide as an oil. IR includes peaks at 1660cm⁻¹

(CO) and 3300-3420cm⁻¹ (NH₂).

From Intermediate 34(c).

5 (d) 2-(4-Aminophenylthio)-N-[(3,4-dimethoxyphenyl)methyl]-N-methyl acetamide as an oil. IR includes peaks at 1645 cm⁻¹ (CO) and 3350cm⁻¹ (NH₂).

From Intermediate 34(d).

10 (e) 2-(4-Aminophenoxy)-N-[2-(4-methoxyphenyl)ethyl]-N-methylacetamide as an oil. IR includes peaks at 1630cm⁻¹ (CO) and 3350-3420cm⁻¹ (NH₂).

From Intermediate 34(e).

15 (f) 2-(4-Aminophenoxy)-N-[(4-methoxyphenyl)methyl]-N-methylacetamide as an oil. IR includes peaks at 1650cm⁻¹ (CO) and 3340-3430cm⁻¹ (NH₂).

From Intermediate 34(f).

20 (g) 2-(4-Aminophenoxy)-N-methyl-N-[(4-methylphenyl)methyl]acetamide as an oil. IR includes peaks at 1650cm⁻¹ (CO) and 3350-3420cm⁻¹ (NH₂).

From Intermediate 34(g).

25 (h) 2-(4-Aminophenoxy)-N-methyl-N-[(4-(methylthio)phenyl)methyl] acetamide as an oil. IR includes peaks at 1660cm⁻¹ (CO) and 3340-3420cm⁻¹ (NH₂).

From Intermediate 34(h).

30 (i) 2-(4-Aminophenoxy)-N-ethyl-N-(phenylmethyl)acetamide as an oil. IR includes peaks at 1650cm⁻¹ (CO) and 3350-3430cm⁻¹ (NH₂).

From Intermediate 34(i).

35 Intermediate 36

(a) N-[2-(4-Aminophenoxy)ethyl]-3,4-dimethoxy-N-methylbenzenethanamine

A solution of Intermediate 35(a) (20 g) in THF (200 ml) was added dropwise to a stirred suspension of lithium aluminium hydride in THF (100 ml) at room temperature and the mixture was heated under reflux for 3h. Water was added carefully to the cooled mixture which was then filtered, washed with THF, evaporated and extracted with ether. The combined ethereal extracts were dried and evaporated to give the title compound (11 g) as an oil. IR includes a peak at 3350-3430cm⁻¹ (NH₂).

The following compounds were prepared in a similar manner to Intermediate 36(a):

45 (b) N-[2-(4-Aminophenoxy)ethyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil. IR includes a peak at 3360-3420cm⁻¹ (NH₂).

From Intermediate 35(b).

50 (c) N-[2-(4-Aminophenoxy)ethyl]-N-methylbenzenemethanamine as an oil. IR includes a peak at 3330-3420cm⁻¹ (NH₂).

From Intermediate 35(c).

55 (d) N-[2-(4-Aminophenylthio)ethyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil. NMR includes signals at δ 2.30 (3H,s,N-CH₃) and 3.85 (6H,s,OCH₃).

From Intermediate 35(d).

(e) N-[2-(4-Aminophenoxy)ethyl]-4-methoxy-N-methylbenzenethanamine as an oil. IR includes a peak at 3340-3430cm⁻¹ (NH₂).

5

From Intermediate 35(e).

(f) N-[2-(4-Aminophenoxy)ethyl]-4-methoxy-N-methylbenzenemethanamine as an oil. IR includes a peak at 3350-3430cm⁻¹ (NH₂).

10

From Intermediate 35(f).

(g) N-[2-(4-Aminophenoxy)ethyl]-4-methyl-N-methylbenzenemethanamine as an oil. IR includes a peak at 3350-3430cm⁻¹ (NH₂).

15

From Intermeiate 35(g).

(h) N-[2-(4-Aminophenoxy)ethyl]-N-methyl-4-(methylthio) benzenemethanamine as an oil. IR includes a peak at 3350-3420cm⁻¹ (NH₂).

20

From Intermediate 35(h).

(i) N-[2-(4-Aminophenoxy)ethyl]-N-ethylbenzenemethanamine as an oil. IR includes a peak at 3360-3430cm⁻¹ (NH₂).

25

From Intermediate 35(i).

Intermediate 37

30 (a) 3,4-Dimethoxy-N-methyl-N-[3-(4-nitrophenoxy)propyl] benzeneethanamine

A mixture of 1-(3-bromopropoxy)-4-nitrobenzene (18.7 g) and Intermediate 20(a) (14.1g) were heated for 30 min at 140° and then diluted with water. The mixture was extracted with dichloromethane, and the organic phase was washed with water, dried and concentrated. The residue was purified by column chromatography eluting with dichloromethane/methanol (95:5) to give the title compound (18g) as an oil. NMR includes a signal at d 2.38 (3H,s,N-CH₃).

The following compounds were prepared in a similar manner to Intermediate 37(a):

40 (b) 4-Methoxy-N-methyl-N-[3-(4-nitrophenoxy)propyl] benzeneethanamine as an oil. NMR includes a signal at d 2.40 (3H,s,N-CH₃).

From 1-(3-bromopropoxy)-4-nitrobenzene and Intermediate 20(e).

45 (c) 3,4-Dimethoxy-N-methyl-N-[3-(4-nitrophenoxy)propyl] benzenemethanamine as an oil. NMR includes a signal at d 2.40 (3H,s,N-CH₃).

From 1-(3-bromopropoxy)-4-nitrobenzene and Intermediate 20(b).

50 (d) 3,4-Dimethoxy-N-methyl-N-[3-[(4-nitrophenoxy)propyl]thio]benzenemethanamine as an oil. NMR includes a signal at d 2.40 (3H,s,N-CH₃).

From i-[(3-bromopropoxy)thio]-4-nitrobenzene and Intermediate 20(b).

Intermediate 38

55

(a) N-[3-(4-Aminophenoxy)propyl]-3,4-dimethoxy-N-methylbenzenethanamine

A solution of Intermediate 37(a) (18g) in ethanol (200 ml) was hydrogenated at room temperature in the

presence of 10% palladium on carbon (1 g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (15g) as an oil. IR includes a peak at 3300-3370cm⁻¹ (NH₂).

The following compounds were prepared in a similar manner to Intermediate 38(a):

6 (b) N-[3-(4-Aminophenoxy)propyl]-4-methoxy-N-methylbenzeneethanamine as an oil. IR includes a peak at 3350-3430cm⁻¹ (NH₂).

10 From Intermediate 37(b).

10 (c) N-[3-(4-Aminophenoxy)propyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil. IR includes a peak at 3360-3430cm⁻¹ (NH₂).

15 From Intermediate 37(c).

15 (d) N-[3-[(4-Aminophenyl)thio]propyl]-3,4-dimethoxy-N-methylbenzenemethanamine as an oil. IR includes a peak at 3370-3450cm⁻¹ (NH₂).

20 From Intermediate 37(d).

20 Intermediate 39

9,10-Dihydro-2-(methylthio)-9-oxo-4-acridinecarboxylic acid

25 (i) 2-[(2-Carboxyphenyl)amino]-5-(methylthio)benzoic acid

A mixture of 2-chloro-5-(methylthio)benzoic acid (10 g), anthranilic acid (7 g), potassium carbonate (14 g) and copper (1 g) in 2-(2-methoxyethoxy)ethanol(100 ml) was heated at 180° for 24h. Water (400 ml) was then added, and the catalyst was filtered off. The filtrate was acidified with dilute hydrochloric acid. The 30 resulting precipitate was filtered off, washed with water, dried, and crystallised from methanol to give the title compound (4.5g) as crystals. IR includes peaks at 3300cm⁻¹ (NH) and 1700cm⁻¹ (CO₂H).

(ii) 9,10-Dihydro-2-(methylthio)-9-oxo-4-acridinecarboxylic acid

35 The product of part (i) above (2g) in phosphorus oxychloride (6 ml) was heated at reflux for 1h. The solution was then cooled (to 0°), and water (15 ml) was added slowly. The mixture was then heated at 100° for 10 min and then poured onto cracked ice. The resulting precipitate was filtered off, washed with water, and crystallised from methanol to give the title compound (1.6g). IR includes peaks at 1690cm⁻¹ (CO₂H) and 1620cm⁻¹ (CO).

40 Intermediate 40

N-[4-(3-Bromopropoxy)phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

45 (i) N-[4-(3-Bromopropoxy)phenyl]acetamide

A mixture of N-(4-hydroxyphenyl)acetamide (10 g) and potassium carbonate (11 g) in DMF (200 ml) was stirred for 20 min at room temperature. 1,3-Dibromopropane (35 ml) was then added and stirring was continued for 4 h. The mixture was filtered and the filtrate was concentrated in vacuo. The residue was 50 treated with water and extracted with dichloromethane. The organic phase was washed first with dilute sodium hydroxide, then with water, dried and concentrated to give a solid which was triturated with hexane to give the title compound (14g), MP: 120°.

(ii) 4-(3-Bromopropoxy)benzenamine

55 A mixture of the product of part (i) above (13g) and 5N hydrochloric acid (200 ml) was heated under reflux for 2 h. After cooling, the mixture was basified with sodium hydroxide solution and extracted with dichloromethane. The organic phase was evaporated to give the title compound (7g) as an oil. IR includes a

peak at 3360-3450cm⁻¹ (NH).

(iii) N-[4-(3-Bromopropoxy)phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 A mixture of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1.5 g) and 1-hydroxybenzotriazole (1.1 g) in DMF (50 ml) was stirred at room temperature for 10 min. The product of part (ii) above (1.5g) was then added followed by dicyclohexylcarbodiimide (1.3 g), and the mixture was stirred at room temperature for 16 h and then filtered. The filtrate was concentrated in vacuo, treated with water and extracted with dichloromethane. The combined, dried organic extracts were concentrated to give the title compound (0.5g)

10 which was recrystallised from acetonitrile, MP 126°.

Intermediate 41

N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-nitrophenylaminocarbonylmethanamine

15 A mixture of Intermediate 20(b) (2.8g), Intermediate 56 (3g) and potassium carbonate (2.3g) in DMF (50ml) was heated at 60° for 24h. The mixture was then evaporated, extracted with dichloromethane, washed with water, dried and concentrated to give a solid which was recrystallised from diethyl ether to provide the title compound (3.7g). MP: 120°.

20 Intermediate 42

N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-4-aminophenylaminocarbonylmethanamine

25 A solution of Intermediate 41 (3.6g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium on carbon (500mg). After hydrogen absorption was completed the catalyst was removed by filtration and the filtrate was concentrated to give the title compound (3.5g).
NMR includes signals at d 2.5 (3H,s,N-CH₃); 3.8 (6H,s,OCH₃).

30 Intermediate 43

N-[2-(4-Aminophenylamino)ethyl]-3,4-dimethoxy-N-methylbenzenemethanamine

35 A solution of Intermediate 42 (3.5g) in THF (50ml) was added dropwise to a stirred suspension of lithium aluminium hydride in THF (30ml) at room temperature and the mixture was heated under reflux for 48h. Water was added carefully to the cooled mixture which was then filtered on a celite pad. The filtrate was evaporated to dryness and upon column chromatography (dichloromethane-methanol), the remaining residue gave the title compound (1.4g).

NMR includes signals at d 2.15 (3H,s,N-CH₃); 2.5 and 3 (4H,2t,-CH₂-CH₂); 3.7 (6H,s,OCH₃).

40 Intermediate 44

9,10-Dihydro-5,7-dimethoxy-9-oxo-4-acridinecarboxylic acid

45 A mixture of 2-iodoisophthalic acid (5.8g), 2,4-dimethoxy-aniline (4.3g) and cuprous chloride (1g) in 2,3-butanediol (20ml) and toluene (10ml) was heated to 120°. After most of toluene has distilled off, N-ethylmorpholine (10ml) was added and the mixture was stirred at 120° for one hour. After cooling and dilution with 2N potassium carbonate the solution was filtered on celite. The filtrate was acidified with 2N hydrochloric acid and the greenish precipitate was recovered by filtration.

50 The product (4g) was heated in polyphosphoric acid (50g) at 120° for 1.5 hour to give the title compound which was recovered as a solid (1.5g) by precipitation with water and purified by dissolving in 1N sodium hydroxide and reprecipitation with acetic acid (pH 4).

55

Analysis Found :	C,62.1;	H,4.6;	N,4.3;
C ₁₆ H ₁₃ NO ₅ , 0.5 H ₂ O Requires :	C,62.3;	H,4.6;	N,4.5%.

The following acid was prepared in a similar manner to Intermediate 44.

Intermediate 45

9,10-Dihydro-6,7,8-trimethoxy-9-oxo-4-acridinecarboxylic acid (1.5g). IR includes a peak at 1620cm^{-1} (CO).

5 From 3,4,5-trimethoxyaniline (3.8g) and 2-iodoisophthalic acid (5g).

Intermediate 463-(2-Bromoethyl)nitrobenzene

10 Phosphorus tribromide (0.94ml) was added dropwise to a solution of 3-nitrophenethyl alcohol (5g) in anhydrous diethyl ether (30ml) at 0° . The mixture was stirred at room temperature for 2 hours and then treated with a solution of potassium carbonate and then water. The organic layer was dried and concentrated in vacuo to give the title compound as an oil (4.51g).

15 NMR includes d 3.25 (m,2H,CH₂-Ph) and 3.55 (m,2H,CH₂-Br).

Intermediate 47(a) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-nitrobenzenethanamine

20 A mixture of Intermediate 46 (2.2g), Intermediate 20(b) (1.71g) and potassium carbonate (1.58g) in DMF (50ml) was heated at 60° for 36 hours. The mixture was filtered and the filtrate concentrated in vacuo. The residue was treated with water and extracted with methylene chloride. The organic extract was dried, concentrated and purified by column chromatography on silica gel eluting with methylene chloride/methanol (99:1) to give the title compound as an oil (1g).

25 NMR includes d 2.2 (s,3H,N-CH₃) and 3.7 (s,6H,2xOCH₃).

In the same way was prepared the following compound:

(b) N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-(3-nitrophenoxy) propanamine

30 From 3-(3-bromopropoxy)nitrobenzene and Intermediate 20(b).

NMR includes d 2.2 (s,3H,N-CH₃), 3.35 (s,2H,N-CH₂-Ph) and 3.8 (s,6H,2xOCH₃).

Intermediate 48(a) 3-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenethanamine

35 A solution of Intermediate 47(a) (1g) in ethanol (50ml) was hydrogenated at room temperature in presence of 10% palladium-on-carbon (0.15g). After the hydrogen absorption was completed, the catalyst

40 was filtered off and the solution concentrated to give the title compound as an oil (0.8g).

NMR includes d 2.25 (s,3H,N-CH₃), 3.4 (s,2H,NH₂) and 3.8 (s,6H,2xOCH₃).

In the same way was prepared the following compound:

(b) N-[3-(3-Aminophenoxy)propyl]-3,4-dimethoxy-N-methylbenzenemethanamine

45 From Intermediate 47(b).

NMR includes d 2.2 (s,3H,N-CH₃), 2.7 (s,2H,NH₂), 3.4 (s,2H,N-CH₂-Ph) and 3.7 (s,6H,2xOCH₃).

Intermediate 49N-[(3,4-Dimethoxyphenyl)methyl]-N-methyl-3-(3-nitrophenyl)-2-propenamide

50 A mixture of 3-nitrocinnamic acid (10g) and 1-hydroxybenzotriazole (8.26g) in DMF (100ml) was stirred at room temperature for 10 minutes. Intermediate 20(b) (9.2g) was added followed by dicyclohexylcarbodiimide (10.63g). The mixture was stirred at room temperature for 16 hours and then filtered. The filtrate was concentrated in vacuo, treated with dilute hydrochloric acid solution, then dilute sodium hydroxide solution and extracted with methylene chloride. The organic extract was dried and concentrated to give the title compound (15.63g).

NMR includes d 3.1 (s,3H,N-CH₃) and 3.75 (s,6H,2xOCH₃).

Intermediate 50

5 3-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamide

A solution of Intermediate 49 (10g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (1g). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate concentrated *in vacuo*. The residue was purified by column chromatography on 10 silica gel eluting with methylene chloride/methanol (98:2) to give the title compound as an oil (5.56g). NMR d 2.7 (s,2H,N-CH₃) and 3.85 (s,6H,2xOCH₃).

Intermediate 51

15 3-Amino-N-[(3,4-dimethoxyphenyl)methyl]-N-methylbenzenepropanamine

A solution of Intermediate 50 (5g) in THF (100ml) was added dropwise to a stirred suspension of lithium aluminium hydride (2.31g) in THF (80ml) at room temperature and the mixture was heated under reflux for 2 hours. Water (20ml) was carefully added to the cooled mixture which was then filtered. The filtrate was 20 concentrated, treated with water and extracted with diethyl ether. The organic extract was dried, evaporated and the product purified by column chromatography on silica gel eluting with methylene chloride/methanol (97:3) to give the title compound as an oil (2.46g).

NMR includes d 2.1 (s,3H,N-CH₃), 3.35 (s,2H,N-CH₂-Ph) and 3.7 (s,6H,2xOCH₃).

25 Intermediate 52

4-(3-Methoxy-4-nitrophenyl)-3-buten-1-ol

The Wittig reaction in THF (100ml) between 3-methoxy-4-nitrobenzaldehyde (1) (2g) and 3-hydroxy-30 propyltriphenylphosphonium bromide (2) (5.3g) in presence of a solution of n-butyllithium (1.6M) in hexane (16.5ml) gave the title compound (2.6g) as an oil.

NMR includes signal at d 3.4(2H,t,CH₂OH); 3.6(3H,s,OCH₃).

(1) CA113 (19): 171567 w

(2) A.R. Hands and A.J.H. Mercer, J. Chem. Soc. (c), (1968) 2448.

35

Intermediate 53

4-(4-Bromo-1-butenyl)-2-methoxy-1-nitrobenzene

40 Phosphorus tribromide (0.33ml) was added dropwise to a solution of Intermediate 52 (2.6g) in anhydrous diethyl ether (10ml) at 0°. The mixture was stirred at room temperature for 1 hour, then washed with a solution of potassium carbonate (1M) and with water. The organic layer was dried and concentrated *in vacuo* to give the title compound (3.3g) as a yellow oil. NMR includes signals at d 3.35(2H,t,CH₂-Br); 3.8-(3H,s,O-CH₃).

45

Intermediate 54

N-[4-(3-Methoxy-4-nitrophenyl)-3-butenyl]-3,4-dimethoxy-N-methylbenzenemethanamine

50 A mixture of Intermediate 53 (3.3g), Intermediate 20(b) (2.5g) and potassium carbonate (1.9g) in DMF (20ml) was stirred at room temperature for 48h. The mixture was filtered and the filtrate was evaporated. The residue was taken into water and extracted with dichloromethane. The organic layer was washed with water, dried, filtered and evaporated. The oily residue was then purified by silica gel column chromatography eluting with dichloromethane/methanol (95:5) to give the title compound (3.4g) as an oil. NMR 55 includes signals at d 2.1(3H,s,N-CH₃); 3.7(6H,s,2xOCH₃); 3.8(3H,s,OCH₃).

Intermediate 55

4-Amino-N-[(3,4-dimethoxyphenyl)methyl]-3-methoxy-N-methylbenzenebutanamine

5 A solution of Intermediate 54 (1.2g) in a mixture of ethanol (50ml) and ethyl acetate (20ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.1g). After the hydrogen absorption was completed, the catalyst was filtered off and the solution concentrated to give the title compound (1g) as an oil. NMR includes signals at d 2.1 (3H,s,N-CH₃); 3.65(3H,s,O-CH₃); 3.7(6H,s,2xOCH₃).

Intermediate 5610 2-Chloro-N-(4-nitrophenyl)acetamide

15 Chloroacetyl chloride (11ml) was added dropwise to a stirred mixture of potassium carbonate (18.8g) and 4-nitroaniline (15g) in DMF (100ml) maintained at 0°. The mixture was then allowed to stand overnight at room temperature and poured into crushed ice. A yellow solid was recovered and crystallised from toluene containing isopropyl alcohol (10%) to give the title compound (10g), MP: 180°.
NMR includes signals at d 4.1(2H,s,COCH₂Cl); 7.4-8.1(4H,m,aromatics); 10.3(1H,bs,NH).

Intermediate 5720 3,4-Dihydro-6,7-dimethoxy-N-(4-nitrophenyl)-2(1H)-isoquinolineacetamide

25 A mixture of Intermediate 56 (10.3g), potassium carbonate (8g) and 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline (9.3g) in DMF (100ml) was heated overnight at 60°. After cooling, the reaction mixture was poured onto ice and the insoluble material recovered and dried to give the title compound, MP: 173-178°.
NMR includes signals at d 2.8(4H,s,2xCH₂); 3.2(2H,s,COCH₂-N); 3.7(2H,s,N-CH₂-Ph); 3.7(6H,m,2xOCH₃); 6.2-8.15(6H,m,aromatics); 9.3(1H,bs,NHCO).

Intermediate 5830 N-(4-Aminophenyl)-3,4-dihydro-6,7-dimethoxy-2(1H)-isoquinolineacetamide

35 A suspension of Intermediate 57 (15g) and 10% palladium-on-carbon (1g) in ethanol (200ml) was stirred at room temperature under a slight overpressure of hydrogen. After 2h the catalyst was filtered off, and washed with dichloromethane/methanol (9:1). The filtrate and washing were concentrated and the crystalline residue gave upon washing with ethanol and drying the title compound (10.6g), MP: 185°. NMR includes signals at d 2.8(4H,s,2xCH₂); 3.15(2H,s,CO-CH₂-N); 3.6(2H,s,Ph-CH₂-N); 3.7(6H,s,2xOCH₃); 6.15-7.3-(6H,m,aromatics); 8.65(1H,bs,CONH).

Intermediate 5940 N-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]-1,4-benzenediamine

45 A solution of borane in tetrahydrofuran (1M; 35.4ml) was added to a stirred solution of Intermediate 58 (2g) in THF (150ml). After 4h of refluxing the reaction mixture was cooled, treated with concentrated hydrochloric acid to make the solution up to 3N in hydrochloric acid and then refluxed again for 15 min.
10N Sodium hydroxide was added and the mixture was extracted with dichloromethane. The organic layer was washed with water, dried and concentrated to give a residue which after purification by silica gel column chromatography eluting with toluene/isopropylamine (95:5) gave the title compound as an oil (1.2g).
NMR includes signals at d 2.6(4H,bs,Ph-CH₂-CH₂-N); 3.45(4H,s,CH₂-NHPh and PhCH₂-N); 3.6-(6H,s,2xOCH₃); 6.3(6H,s,aromatics).

Intermediate 604-[2-(2,3-Dihydro-5,6-dimethoxy-1H-isoindol-2-yl)ethyl]benzenamine

55 4,5-Bischloromethyl veratrol (2.35g: S. H. Wood, M. A. Peny and C. C. Tung, J. A. C. S., (1950), 72, 2989-2991) was added at room temperature to a stirred suspension of 50% aqueous sodium hydroxide (5 ml), toluene (25 ml), 4-aminophenylethylamine (1.5g) and Aliquat (0.2g). The heterogeneous mixture was

stirred at room temperature for 16 hours, poured in water and extracted with methylene chloride. The organic layer was dried and the solvent evaporated in vacuo. The residue was purified by column chromatography on silica gel eluting with methylene dichloride/methanol (95:5) to give the title compound as a solid (0.8g), MP: 150°. NMR includes signals at d 2.7(4H,m,Ph-CH₂-CH₂-N); 4.6(2H,bs,NH₂); 3.7-6(6H,s,2xOCH₃); 3.8(4H,s,2xN-CH₂Ph); 6.2-7.0(6H,m,aromatics).

Intermediate 61

1-(4-Nitrophenyl)-2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethanone hydrobromide

10 A solution of 6,7-dimethoxy-1,2,3,4-tetrahydroisoquinoline (15.63g) and 2-bromo-4'-nitroacetophenone (18.47g) in a mixture of ethanol (150ml) and methylene chloride (150ml) was heated at 60° for 24 hours. After cooling to room temperature yellow crystals appeared. These were collected by filtration and dried in vacuo to give the title compound (9.4g); MP: 216°. NMR(D₆-DMSO) includes signals at d 3.6(6H,s,2xOCH₃); 4.2(2H,s,N-CH₂-Ph); 4.95(2H,s,CO-CH₂-N); 6.6(2H,aromatics isoquinoline); 8(4H,m,aromatics).

Intermediate 62

3,4-Dihydro-6,7-dimethoxy-a-(4-nitrophenyl)-2(1H)-isoquinolineethanol

20 To a suspension of Intermediate 61 (9.4g) in methanol (600ml) was added portionwise sodium borohydride (2.44g) and the mixture was stirred at room temperature for 16 hours. The reaction was diluted with water (200ml), filtered and evaporated in vacuo. The residue was extracted with methylene chloride and washed with water. The organic layer was dried and evaporated in vacuo to give the title compound (1.15g), 25 after crystallisation from ethanol, MP: 130°. NMR includes signals at d 2.4-3.1(6H,m,3xCH₂); 3.7-(6H,s,2xOCH₃); 4.2(1H,bs,OH); 4.8(1H,m,H-C-OH); 6.1-8.1(6H,m,aromatics).

Intermediate 63

a-(4-Aminophenyl)-3,4-dihydro-6,7-dimethoxy-2(1H)-isoquinolineethanol

30 A solution of Intermediate 62 (2.4g) in ethanol (200ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.3g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (1.9g) as a white solid, MP: 168°. 35 NMR includes signals at d 2.4-2.9(6H,m,3xCH₂); 3.5(2H,bs,NH₂); 3.7(6H,s,2xOCH₃); 4.55(1H,t,H-COH); 6.25-7.1(6H,m,aromatics).

Intermediate 64

2-Bromo-N-methyl-N-[(4-nitrophenyl)methyl]acetamide

40 To a solution of bromoacetyl bromide (30g) in methylene chloride (20ml) at 0° was added a solution of N-methyl-4-nitrobenzenemethanamine (8.3g; G. I. Wilson, J. Chem. Soc., 1926, 2461) in methylene chloride (10ml) and triethylamine (12ml). The reaction was stirred 5 min. at 0° and then water (20ml) was added. The methylene chloride layer was dried and evaporated in vacuo. The residue was purified by column chromatography eluting with methylene chloride/methanol (97:3) to give the title compound (15g) as an oil. NMR includes signals at d 3.1(3H,s,N-CH₃); 3.9(2H,s,CH₂Br); 4.55(2H,s,Ph-CH₂-N); 7.0-8.3(4H,m,aromatics).

Intermediate 65

3,4-Dihydro-6,7-dimethoxy-N-methyl-N-[(4-nitrophenyl)methyl]-2(1H)-isoquinolineacetamide

50 A mixture of Intermediate 64 (1.8g), 6,7-dimethoxy-1,2,3,4-tetrahydroisoquinoline (1.4g) and potassium carbonate (1.6g) in DMF (150ml) was stirred overnight. After removal of insoluble material by filtration the solvent was evaporated in vacuo and the residue partitioned between dichloromethane and water. The organic phase was dried, then concentrated under reduced pressure and the product, after purification by column chromatography eluting with methylene chloride/methanol (96:4), gave the title compound (1.65g). NMR includes signals at d 2.8(4H,m,2xCH₂); 3.0(3H,s,N-CH₃); 3.33(2H,s,CO-CH₂-N); 3.6(2H,s,N-CH₂-Ph);

3.7(6H,s,2xOCH₃); 4.55(2H,s,Ph-CH₂-NHCO); 6.2-8.1(6H,m,aromatics).

Intermediate 66

5 N-[(4-Aminophenyl)methyl]-3,4-dihydro-6,7-dimethoxy-N-methyl-2(1H)-isoquinolineacetamide

A solution of Intermediate 65 (1.65g) in ethyl acetate (100ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium-on-carbon (0.34g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound - (1.43g) as a white solid, MP: 175-215°. NMR includes signals at d 2.8(7H,m,NCH₃ and 2xCH₂); 3.2-(2H,s,CO-CH₂-N); 3.5(2H,s,N-CH₂-Ph); 3.7(6H,s,2xCH₃).

Intermediate 67

15 N-[(4-Aminophenyl)methyl]-3,4-dihydro-6,7-dimethoxy-N-methyl-2(1H)-isoquinolineethanamine

A solution of Intermediate 66 (1.49g) in THF (150ml) was added dropwise to a stirred suspension of lithium aluminium hydride (0.47g) in THF (100ml) at room temperature for 4 hours. Water (5ml) was added carefully to the cooled mixture which was filtered and the filtrate concentrated and the residue extracted with methylene chloride. The organic layer was dried and evaporated. The resulting product was purified by column chromatography on silica gel eluting with methylene chloride/isopropylamine (92:8) to give the title compound as an oil (0.7g). NMR includes signals at d 2.15(3H,s,N-CH₃); 2.55(8H,m,4xCH₂); 3.55(2H,s,NH₂); 3.65(6H,s,2xOCH₃); 6.3-7.1(6H,m,aromatics)

25 Intermediate 68

2-[[(3,4-Dimethoxyphenyl)methyl]methylamino]-N-methyl-N-[(4-nitrophenyl)methyl]acetamide

A mixture of Intermediate 64 (4.3g), Intermediate 20(b) (3.26g) and potassium carbonate (4.14g) in DMF (100ml) was stirred overnight. The mixture was filtered, and the filtrate concentrated in vacuo to a residue which was extracted with methylene chloride. After washing with water and drying, the organic layer was evaporated to a syrup which was purified by column chromatography on silica gel eluting with ethyl acetate/cyclohexane (1:1) to give the title compound as an oil (5.7g). NMR includes signals at d 2.3(3H,s,N-CH₃); 3.7(6H,s,2xOCH₃); 4.5(2H,s,Ph-CH₂-NHCO).

35 Intermediate 69

N-[(4-Aminophenyl)methyl]-2-[[(3,4-dimethoxyphenyl)methyl]-methylamino]-N-methylacetamide

40 A solution of Intermediate 68 (5.7g) in a mixture of ethyl acetate/methanol (1:2) (100ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium-on-carbon (0.8g). After hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated to give the title compound (5.2g) as an oil. NMR includes signals at d 3.8(6H,s,2xOCH₃); 4.5(2H,s,Ph-CH₂-NCO).

45 Intermediate 70

N-[(4-Aminophenyl)methyl]-N'-[(3,4-dimethoxyphenyl)methyl]-N,N'-dimethyl-1,2-ethanediamine

50 A solution of Intermediate 69 (5.2g) in THF (150ml) was added dropwise at room temperature to a stirred suspension of lithium aluminium hydride (1g) in THF (50ml). After 4 hours, water (10ml) was added carefully to the cooled mixture which was then filtered. The filtrate was concentrated to dryness and the residue diluted with methylene chloride and extracted with hydrochloric acid (1M). The aqueous layer was basified with an aqueous solution of sodium hydroxide (1M) and extracted with methylene chloride. The organic layer was dried and then concentrated in vacuo. The residue was purified by column chromatography on silica gel eluting with cyclohexane/methylene chloride/isopropylamine (5:4:1) to give the title compound as an oil (2g). NMR includes signals at d 2.1(6H,s,2xNCH₃); 2.4(4H,s,2xNCH₂); 3.2(4H,m,2xN-CH₂-Ph); 3.6(6H,s,2xOCH₃); 3.85(2H,s,NH₂); 6.1-7.5(7H,m,aromatics).

Intermediate 713,4-Dimethoxy-N-methyl-N-[4-(4-nitrophenyl)-2-butenyl]benzenemethanamine

5 A mixture of Intermediate 20(b) (9g), potassium carbonate (8g) and 1-chloro-4-(4-nitrophenyl)-2-butene (10.6g; Morgan and al., J. Med. Chem., 8, (1965), 1398-1405) in 4-methyl-2-pentanone (300ml) was refluxed for 18 hours. After cooling, the mixture was filtered and evaporated in *vacuo*. The residue was purified by column chromatography eluting with methylene chloride/methanol (97.5:2.5) to give the title compound (2g) as an oil. NMR includes signals at d 2.2(3H,s,N-CH₃); 3.9(6H,s,2xOMe); 5.7(2H,m,double bond); 6.9-10 (3H,m,aromatics Ph(OMe)₂); 7.4 and 8.15(4H,2d,aromatics PhNO₂).

Intermediate 72N-[4-(4-Aminophenyl)-2-butenyl]-3,4-dimethoxy-N-methylbenzenemethanamine

15 Intermediate 71 (1.7g) was dissolved at room temperature with stirring in a mixture of methanol (50ml) and concentrated hydrochloric acid (2ml). Iron powder (1.5g) was then added slowly, and the reaction mixture was heated under reflux for 1h. The mixture was then evaporated, basified with sodium hydroxide and extracted with diethyl ether. The organic layer was dried and evaporated in *vacuo* to give the title compound (0.21g) as an oil. NMR includes signals at d 2.15(3H,s,N-CH₃); 3.8(6H,s,2xOMe); 5.55-(2H,m,double bond); 6.3-7.2(7H,m,aromatics).

Intermediate 733,4-Dimethoxy-N-methyl-N-[3-(4-nitrophenyl)-2-propenyl] benzenemethanamine

25 A mixture of Intermediate 20(b) (3.6g), 1-chloro-3-(4-nitrophenyl)-2-propene (4.8g; Cignarella and al., J. Med. Chem., 8, (1965), 326-329) and potassium carbonate (3.5g) in 4-methyl-2-pentanone (60ml) was refluxed for 3 hours. After cooling, the mixture was filtered and the filtrate was evaporated in *vacuo*. The residue was purified by column chromatography eluting with methylene chloride/methanol (95:5) to give the title compound (4.9g) as an oil. NMR includes signals at d 2.25(3H,s,NCH₃); 3.2(2H,d,N-CH₂-CH=CH); 3.5-(2H,s,NCH₂Ph); 3.85(6H,s,2xOMe); 6.55(2H,m,double bond); 6.8(3H,d,aromatics Ph(OMe)₂); 7.4 and 8.1-(4H,2d,aromatics PhNO₂).

Intermediate 744-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]-1-propenyl] benzenamine

35 Intermediate 73 (4.8g) was dissolved in a mixture of methanol (100ml) and concentrated hydrochloric acid (10ml) at room temperature with stirring. Iron powder (5g) was then added slowly and the reaction mixture was refluxed for 0.5h. After cooling, the mixture was evaporated, diluted with water (20ml), basified with sodium hydroxide solution, concentrated and extracted with diethyl ether. The organic layer was dried and evaporated to give the title compound (3.95g) as an oil. NMR includes signals at d 2.2(3H,s,NCH₃); 3.15(2H,d,N-CH₂-CH=CH); 3.5(2H,s,NCH₂Ph); 3.6(2H,s,NH₂); 3.8(6H,s,2xOMe); 5.7-7.6(9H,m,aromatics and double bond).

Intermediate 751,2,3,4-Tetrahydro-6-methoxy-2-[2-(4-nitrophenyl)ethyl]isoquinoline

50 A mixture of 1-(2-bromoethyl)-4-nitrobenzene (6.4g), 1,2,3,4-tetrahydro-6-methoxyisoquinoline (4.6g; Daniel J. Sall and Gary L. Grunewald, J. Med. Chem. 1987, 30, 2208-2216) and potassium carbonate (9.7g) in DMF (150ml) was stirred at 50° for 15 h. The mixture was evaporated to dryness and the residue was extracted with dichloromethane. The organic layer was washed with water, dried, filtered and evaporated. 55 The residue was then purified by column chromatography eluting with dichloromethane/methanol (98:2) to give the title compound (2g) as an oil which solidified on standing. NMR includes signals at d 3.6(2H,m,N-CH₂Ar), 3.7 (3H,s,OCH₃).

Intermediate 764-[2-(1,2,3,4-Tetrahydro-6-methoxy-2-isoquinoliny)ethyl]-benzenamine

5 A solution of Intermediate 75 (2g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.2g). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated in vacuo to give the title compound (1.8g) as an orange oil which solidified on standing.

NMR includes signals at d 3.4 (2H,s,NH₂), 3.55 (2H,s,N-CH₂Ar), 3.65 (3H,s,OCH₃).

10

Intermediate 771,2,3,4-Tetrahydro-6,7-dimethoxy-2-[3-(3-nitrophenyl)-1-oxo-2-propenyl]isoquinoline

15 A mixture of 3-nitrocinnamic acid (10g) and 1-hydroxybenzotriazole (8.2g) in DMF (100ml) was stirred at room temperature for 10 min. 1,2,3,4-Tetrahydro-6,7-dimethoxy-isoquinoline (10g) was then added, followed by dicyclohexylcarbodiimide (10.6g) and the mixture was stirred at 50° for 48 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide and extracted with dichloromethane. The dried organic extract was evaporated and purified by column chromatography eluting with 20 dichloromethane/methanol (97:3) to give the title compound (7.8g). NMR includes a signal at d 3.85 (6H,s,OCH₃).

Intermediate 782-[3-(3-Aminophenyl)-1-oxopropyl]-1,2,3,4-tetrahydro-6,7-dimethoxy-isoquinoline

A solution of Intermediate 77 (7.8g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (1g). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate concentrated in vacuo to give the title compound (6.8g).

30 IR: Freq CO: 1640cm-1, Freq NH₂: 3450cm-1.

Intermediate 793-[3-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl] benzenamine

35 A solution of Intermediate 78 (6.8g) in THF (100ml) was added dropwise to a stirred suspension of lithium aluminium hydride (3g) in THF (100ml) at room temperature and the mixture was heated under reflux for 3 h. Water was then added carefully to the cooled mixture which was filtered, evaporated and extracted with ether. The extract was dried and evaporated to give the title compound (5.4g) as an oil which solidified 40 on standing.

IR: Freq NH₂: 3350-3450 cm-1.

Intermediate 801-[(3,4-Dimethoxyphenyl)methyl]methylamino]-3-(4-nitrophenoxy)-2-propanol

A mixture of 1,2-epoxy-3-(4-nitrophenoxy)propane (6g; Sigma) and Intermediate 20(b) (5g) in isopropanol (100ml) was heated under reflux for 18 h and evaporated. The oily residue was crystallised from ether to give the title compound (8.3g) as a white solid.

50 NMR includes signals at d 2.3 (3H,s,N-CH₃), 3.9 (6H,s,OCH₃).

Intermediate 811-(4-Aminophenoxy)-3-[(3,4-dimethoxyphenyl)methyl]methylamino]-2-propanol

55 A solution of Intermediate 80 (8g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.8g). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate concentrated in vacuo. The oily product was then purified by column

chromatography eluting with dichloromethane/methanol (95:5) to give the title compound (5.8g) as an oil. NMR includes signals at d 2.25 (3H,s,N-CH₃), 3.8 (6H,s,OCH₃).

Intermediate 82

5

3,4,5-Trimethoxy-N-methyl-N-[3-(4-nitrophenoxy)propyl]benzene methanamine

A mixture of 1-(3-chloropropoxy)-4-nitrobenzene (4.6g), 3,4,5-trimethoxy-N-methylbenzenemethanamine (4.1g; Sigma) and potassium carbonate (2.9g) in DMF (60ml) was heated at 70° for 24 h. The mixture was 10 then filtered and the filtrate evaporated. The residue was taken up in water and extracted with dichloromethane. The organic layer was washed with water, dried, evaporated and purified by column chromatography eluting with dichloromethane/methanol (99:1) to give the title compound (5.8g) as a yellow 15 oil. NMR includes signals at d 2.15 (3H,s,N-CH₃), 3.3 (2H,s,CH₂-Ar), 3.7 (9H,s,OCH₃).

16

Intermediate 83

N-[3-(4-Aminophenoxy)propyl]-3,4,5-trimethoxy-N-methylbenzenemethanamine

A solution of Intermediate 82 (5.8g) in ethanol (100ml) was hydrogenated at room temperature in the 20 presence of 10% palladium-on-carbon (0.5g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (5.1g) as an oil. NMR includes signals at d 2.25 (3H,s,N-CH₃), 3.5 (2H,s,CH₂-Ar), 3.8 (9H,s,OMe).

Intermediate 84

25

1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[(4-methoxy-3-nitrophenyl)acetyl]isoquinoline

A mixture of 4-methoxy-3-nitrophenylacetic acid (1.2g) and 1-hydroxybenzotriazole (0.95g) in DMF (30ml) was stirred at room temperature for 10 min 1,2,3,4-Tetrahydro-6,7-dimethoxy-isoquinoline (1.1g) in 30 DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (1.2g) and the mixture was stirred at room temperature for 6 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide and extracted with ethyl acetate. The dried organic extract was evaporated to give the title compound (1.6g) as an oil which crystallised from ethanol as a white solid, MP 175°. IR: Freq CO: 1650cm⁻¹.

35

Intermediate 85

2-[(3-Amino-4-methoxyphenyl)acetyl]-1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline

A solution of Intermediate 84 (1.6g) in ethanol (50ml) was hydrogenated at room temperature in the 40 presence of 10% palladium-on-carbon (0.3g). After hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (1.4g) as an oil. IR: Freq CO: 1650cm⁻¹, Freq NH₂: 3340-3440 cm⁻¹.

45

Intermediate 86

5-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]-2-methoxybenzenamine

A solution of Intermediate 85 (1.4g) in THF (30ml) was added dropwise to a stirred suspension of 50 lithium aluminium hydride (0.9g) in THF (50ml) at room temperature and the mixture was heated under reflux for 3 h. Water was then added carefully to the cooled mixture which was then filtered, evaporated and extracted with ether. The extract was dried and evaporated to give the title compound (1.2g) as an oil which solidified on standing.

55

IR: Freq NH₂: 3340-3440 cm⁻¹.

Intermediate 87

1,2,3,4-Tetrahydro-2-[3-(4-nitrophenoxy)propyl]isoquinoline

A mixture of 1-(3-bromopropoxy)-4-nitrobenzene (10g), 1,2,3,4-tetrahydroisoquinoline (5.1g) and potassium carbonate (10.6g) in DMF (100ml) was stirred at 70° for 24 h. The mixture was then filtered and the filtrate evaporated. The residue was taken up with water and extracted with dichloromethane. The organic layer was washed with water, dried, evaporated and purified by column chromatography eluting with dichloromethane/methanol (96:4) to give the title compound (8.8g) as a yellow oil. NMR includes signals at 5 d 3.6 (2H,s,N-CH₂Ar), 4.1 (2H,t,O-CH₂).

Intermediate 88

10 4-[3-(1,2,3,4-Tetrahydro-2-isoquinoliny)propoxy]benzenamine

Intermediate 87 (8.8g) was dissolved in a mixture of methanol (80ml) and concentrated hydrochloric acid (50ml) at room temperature with stirring. Iron powder (7.9g) was then added portionwise and the mixture was heated under reflux for 2 h. The mixture was then cooled, poured onto ice, basified with sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water, dried and evaporated to give the title compound (4.5g) as a red oil. NMR includes signals at d 3.7 (2H,s,N-CH₂Ar), 3.9 (2H,t,O-CH₂).

Intermediate 89

20 1,2,3,4-Tetrahydro-7-methoxy-2-[2-(4-nitrophenyl)ethyl]isoquinoline

A mixture of 1-(2-bromoethyl)-4-nitrobenzene (3.7g), 1,2,3,4-tetrahydro-7-methoxyisoquinoline (2.7g; Daniel J. Sall and Gary L. Grunewald, J. Med. Chem. 1987, 30, 2208-2216) and potassium carbonate (6.7g) in isopropanol (150ml) was stirred under reflux for 48 h. The mixture was evaporated to dryness, and the residue was extracted with dichloromethane. The organic layer was washed with water, dried, filtered and evaporated. The residue was then purified by column chromatography eluting with dichloromethane/methanol (99:1) to give the title compound (1.6g) as an orange solid, MP: 92-94°. NMR includes signals at d 3.6 (2H,m,N-CH₂Ar), 3.7 (3H,s,OCH₃).

30 Intermediate 90

4-[2-(1,2,3,4-Tetrahydro-7-methoxy-2-isoquinoliny)ethyl]-benzenamine

35 A solution of Intermediate 89 (1.6g) in ethanol (100ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.16g). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated in vacuo to give the title compound (1.4g) as a white solid, MP: 82-84°.

NMR includes signals at d 3.4 (2H,s,NH₂), 3.45 (2H,s,N-CH₂Ar), 3.55 (3H,s,OCH₃).

40 Intermediate 91

1,2,3,4-Tetrahydro-6,7-dimethoxy-2-[2-(3-nitrophenyl)ethyl] isoquinoline

45 A mixture of 1-(2-bromoethyl)-3-nitrobenzene (2.3g), 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (2.3g) and potassium carbonate (3g) in DMF (50ml) was heated at 50° for 12 h. The mixture was then filtered and the filtrate evaporated. The residue was then taken up in water, extracted with dichloromethane, dried, evaporated and purified by column chromatography eluting with dichloromethane/methanol (99:1) to give the title compound (1.4g) as a yellow oil. NMR includes signals at 50 d 3.6 (2H,s,N-CH₂Ar), 3.75 (6H,s,OCH₃).

Intermediate 92

3-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl] benzenamine

55 A solution of Intermediate 91 (1.4g) in ethanol (50ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.14g). After hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated in vacuo to give the title compound (1.15g) as a yellow oil which

solidified.

NMR includes signals at d 3.6 (2H,s,N-CH₂Ar), 3.75 (6H,s,OCH₃), 4.5 (2H,s,NH₂).

Intermediate 93

5

N-[(3,4-Dimethoxyphenyl)methyl]-4-methoxy-N-methyl-3-nitrobenzeneethanamide

A mixture of 4-methoxy-3-nitrobenzeneacetic acid (1.2g; CA 87, 84684h) and 1-hydroxybenzotriazole (0.95g) in DMF (30ml) was stirred for 10 min. Intermediate 20(b) (1.1g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (1.2g) and the mixture was stirred at room temperature for 6 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide and extracted with ethyl acetate. The dried, organic extract was evaporated to give an oil which was purified by column chromatography eluting with dichloromethane/methanol (95:5) to give the title compound (1.5g) as an oil.

IR: Freq CO: 1640cm-1.

15

Intermediate 94

3-Amino-N-[(3,4-dimethoxyphenyl)methyl]-4-methoxy-N-methylbenzeneacetamide

20 A solution of Intermediate 93 (1.45g) in ethanol (40 ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (0.25g). After the hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (1.2g) as an oil.

IR: Freq CO: 1630cm-1, Freq NH₂: 3350-3450cm-1.

25

Intermediate 95

3-Amino-N-[(3,4-dimethoxyphenyl)methyl]-4-methoxy-N-methylbenzeneethanamine

30 A solution of Intermediate 94 (1.2g) in THF (30ml) was added dropwise to a stirred suspension of lithium aluminium hydride (0.9g) in THF (50ml) at room temperature and the mixture was heated under reflux for 3 h. Water was added carefully to the cooled mixture which was then filtered, washed with THF, evaporated and extracted with ether. The extract was dried and evaporated to give the title compound (1g) as an oil.

IR: Freq NH₂: 3350-3450 cm-1.

35

Intermediate 96

1,2,3,4-Tetrahydro-5,6-dimethoxy-2-[2-(4-nitrophenyl)ethyl] isoquinoline

40 A mixture of 1-(2-bromoethyl)-4-nitrobenzene (0.3g), 1,2,3,4-tetrahydro-5,6-dimethoxyisoquinoline [0.25g; R. D. Haworth, J. Chem. Soc., 2281 (1987); Robin D. Clark, J. Med. Chem., 596-600, 33, (1990)] and potassium carbonate (0.5g) in DMF (25ml) was heated at 60° for 3 h. The mixture was then filtered and the filtrate evaporated. The residue was taken up in water, extracted with dichloromethane, dried, evaporated and purified by column chromatography eluting with dichloromethane/methanol (99:1) to give the title compound (0.3g) as an orange solid, MP:97°. NMR includes signals at d 3.6 (2H,s,N-CH₂Ar), 3.75 (6H,s,OCH₃).

Intermediate 97

50 4-[2-(1,2,3,4-Tetrahydro-5,6-dimethoxy-2-isoquinoliny)ethyl]-benzenamine

A solution of Intermediate 96 (0.3g) in ethanol (20ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (30mg). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated in vacuo to give the title compound (0.22g) as a yellow oil. NMR includes signals at d 3.55 (2H,s,N-CH₂Ar), 3.65-3.85 (8H, OCH₃ and NH₂).

Intermediate 98

1,2,3,4-Tetrahydro-6,7,8-trimethoxy-2-[2-(4-nitrophenyl)ethyl] isoquinoline

5 A mixture of 1-(2-bromoethyl)-4-nitrobenzene (0.34g), 1,2,3,4-tetrahydro-6,7,8-trimethoxyisoquinoline [0.33g; J. Chem. Soc. D, (20), 1296-1297 (1970)] and potassium carbonate (0.5g) in DMF (20ml) was heated at 50° for 12 h. The mixture was then filtered and the filtrate evaporated. The residue was taken up in water, extracted with dichloromethane, dried, evaporated and purified by column chromatography eluting with dichloromethane/methanol (99:1) to give the title compound (0.34g) as a red solid, MP:110°. NMR includes signals at d 3.55 (2H,s,N-CH₂Ar), 3.70 (6H,s,OCH₃), 3.75 (3H,s,OCH₃).

10 Intermediate 994-[2-(1,2,3,4-Tetrahydro-6,7,8-trimethoxy-2-isoquinoliny)ethyl]-benzenamine

15 A solution of Intermediate 98 (0.34g) in ethanol (10ml) was hydrogenated at room temperature in the presence of 10% palladium-on-carbon (50mg). After the hydrogen absorption was completed, the catalyst was filtered off and the filtrate was concentrated in vacuo to give the title compound (0.3g) as a white solid, MP: 92°.

NMR includes signals at d 3.55 (2H,s,N-CH₂Ar), 3.7-3.75 (11H, OCH₃ and NH₂).

20 Intermediate 1001,2,3,4-Tetrahydro-6,7-dimethoxy-2-[2-(4-nitrophenyl)ethyl] isoquinoline

25 A mixture of 1-(2-bromoethyl)-4-nitrobenzene (9.64g), 1,2,3,4-tetrahydro-6,7-dimethoxyisoquinoline hydrochloride (10.59g) and potassium carbonate (17.38g) in isopropanol (150ml) was refluxed for 48h. The mixture was then filtered and the filtrate evaporated to dryness. The resulting residue was taken up in water and extracted with dichloromethane. The organic layer was washed with water, dried and evaporated to give an oil which crystallised in a mixture of 2-propanol and diethyl ether to give the title compound (10.27g). M.p. : 118-119°.

30

Analysis Found :	C,66.48;	H,6.48;	N,8.14;
C ₁₉ H ₂₂ N ₂ O ₄ requires :	C,66.65;	H,6.48;	N,8.18%.

35

Intermediate 1014-[2-(1,2,3,4-Tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl] benzenamine40 Method a :

45 A solution of Intermediate 100 (20g) in ethanol (300ml) was hydrogenated at room temperature and atmospheric pressure in the presence of 10% palladium-on-carbon (2g). After the hydrogen absorption was completed, the catalyst was filtered off and the solution was concentrated to give the title compound (17.2g) as an oil which solidified by scratching in hexane.

Method b :

50 Iron powder (12.44g) was added portionwise at room temperature to a stirred solution of Intermediate 100 (14g) in a mixture of methanol (150ml) and concentrated hydrochloric acid (150ml). After heating under reflux for 45 min, the mixture was cooled, poured onto ice, basified with a solution of sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water, dried and evaporated to give the title compound. M.p.: 128° (ethanol).

55

Analysis Found :	C,72.77;	H,7.80;	N,9.17;
C ₁₉ H ₂₄ N ₂ O ₂ requires :	C,73.05;	H,7.74;	N,8.97%.

Example 19,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide

5

A mixture of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.3g) and 1-hydroxybenzotriazole (0.43g) in DMF (30ml) was stirred at room temperature for 10min. Intermediate 2(c) (1g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (0.66g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The organic layer was then washed with water, dried and evaporated to give a residue which was purified by column chromatography eluting with dichloromethane:methanol (97:3) to give a solid which was recrystallised from isopropanol and filtered off to give the title compound (0.4g), m.p. 215-225°.

15

Analysis Found :	C,72.3;	H,5.9;	N,7.4;
C ₃₄ H ₃₃ N ₃ O ₅ requires :	C,72.5;	H,5.9;	N,7.4%.

20

Example 29,10-Dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide

25

A mixture of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.7g) and 1-hydroxybenzotriazole (0.35g) in DMF (20ml) was stirred at room temperature for 10min. Intermediate 2(b) (0.9g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (0.5g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated to leave an oil which was purified by column chromatography eluting with dichloromethane:methanol (97:3). The resulting solid was recrystallised from acetonitrile and filtered off to give the title compound - (0.26g), m.p. 199°.

35

Analysis Found :	C,67.7;	H,5.9;	N,6.6;	S,5.2;
C ₃₅ H ₃₅ N ₃ O ₅ S(O.5H ₂ O) requires :	C,67.9;	H,5.9;	N,6.8;	S,5.2%.

40

Example 39,10-Dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-4-acridinecarboxamide

45

A mixture of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.5g) and 1-hydroxybenzotriazole (0.5g) in DMF (30ml) was stirred at room temperature for 10min. Intermediate 2(a) (1.27g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (0.76g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated to give a residue which was purified by column chromatography eluting with dichloromethane:methanol (97:3). The solid was recrystallised from isopropanol and filtered off to give the title compound (0.89g), m.p. 190°.

55

Analysis Found :	C,68.6;	H,5.9;	N,6.8;
C ₃₅ H ₃₅ N ₃ O ₆ requires :	C,68.6;	H,6.1;	N,6.9%.

Example 4

5-Fluoro-9,10-dihydro-9-oxo-N-[4-[[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide

5 A mixture of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.5g) in DMF (30ml) was stirred at room temperature for 10min. Intermediate 2(b) (1.4g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (0.8g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated to give a residue which was purified by column chromatography eluting with dichloromethane:methanol (97:3). The 10 solid was recrystallised from isopropanol and filtered off to give the title compound (0.28g), m.p. 162°.

Analysis Found :	C,66.1;	H,5.4;	F,3.0;	N,6.8;	S,5.3;
C ₃₄ H ₃₂ FN ₃ O ₄ S requires :	C,66.3;	H,5.6;	F,3.1;	N,6.8;	S,5.2%.

15

The following compounds were prepared in a similar manner to Examples 1 to 4.

Example 59,10-Dihydro-5-methyl-9-oxo-N-[4-[[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridinecarboxamide

20 The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 2(b) (1.4g) gave, after crystallisation from isopropanol, the title compound (0.45g), m.p. 155°.

25

Analysis Found :	C,68.8;	H,5.9;	N,6.8;	S,5.0;
C ₃₅ H ₃₅ N ₃ O ₄ S(H ₂ O) requires :	C,68.7;	H,6.1;	N,6.8;	S,5.2%.

30

Example 69,10-Dihydro-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-4-acridinecarboxamide

35

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 2(a) (1.1g) gave, after crystallisation from isopropanol, the title compound (0.27g), m.p. 220°.

40

Analysis Found :	C,71.4;	H,5.9;	N,7.3;
C ₃₄ H ₃₃ N ₃ O ₅ (0.5H ₂ O) requires :	C,71.3;	H,6.0;	N,7.3%.

Example 7

45

9,10-Dihydro-9-oxo-N-[4-2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethoxy]phenyl]-4-acridinecarboxamide

50

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.37g) with Intermediate 5(a) (0.51g) gave after crystallisation from isopropanol, the title compound (0.27g), m.p. 154°.

Analysis Found :	C,70.4;	H,5.7;	N,7.5;
C ₃₃ H ₃₁ N ₃ O ₅ (0.5H ₂ O) requires :	C,70.9;	H,5.8;	N,7.5%.

55

Example 8

9,10-Dihydro-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]phenyl]-4-acridincarboxamide

5 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 2(b) (1g) gave, after crystallisation from isopropanol, the title compound (0.04g), m.p. 182°.

Analysis Found :	C,67.3;	H,5.6;	N,6.9;	S,5.25;
C ₃₄ H ₃₃ N ₃ O ₄ S(1.5H ₂ O) requires :	C,67.3;	H,5.9;	N,6.9;	S,5.3%.

10

Example 99,10-Dihydro-5-methyl-9-oxo-N-[4-[4-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)butyl]phenyl]-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 2(d) (1.34g) gave, after crystallisation from ethanol/acetone, the title compound (0.86g), m.p. 140°.

20

Analysis Found :	C,73.1;	H,6.3;	N,6.8;
C ₃₆ H ₃₇ N ₃ O ₄ (H ₂ O) requires :	C,72.8;	H,6.5;	N,7.1%.

25

Example 109,10-Dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-4-acridinecarboxamide

30

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.65g) with Intermediate 5(b) (0.53g) gave, after crystallisation from isopropanol, the title compound (0.3g), m.p. 135°.

35

Analysis Found :	C,70.9;	H,6.0;	N,6.7;
C ₃₅ H ₃₅ N ₃ O ₅ (H ₂ O) requires :	C,70.6;	H,6.3;	N,7.05%.

Example 11

40

9,10-Dihydro-5-methyl-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (0.61g) with Intermediate 5(b) (0.53g) gave, after crystallisation from isopropanol, the title compound (0.45g), m.p. 120°.

45

Analysis Found :	C,73.2;	H,6.15;	N,7.3;
C ₃₅ H ₃₅ N ₃ O ₄ (0.5 H ₂ O) requires :	C,73.7;	H,6.35;	N,7.4%.

50

Example 12

55

5-Fluoro-9,10-dihydro-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 2(c) (0.81 g) gave, after crystallisation from acetonitrile/isopropanol (1:1), the title compound (0.2g), m.p. 212°.

Analysis Found :	C,69.4;	H,5.2;	N,7.8;
C ₃₃ H ₃₀ FN ₃ O ₄ (H ₂ O) requires :	C,69.6;	H,5.6;	N,7.4%.

5

Example 135-Fluoro-9,10-dihydro-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-4-acridinecarboxamide

10

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 5(b) (0.85g) gave, after crystallisation from isopropanol, the title compound (0.4g), m.p. 166°.

15

Analysis Found :	C,70.3;	H,5.4;	N,7.2;
C ₃₄ H ₃₂ FN ₃ O ₄ (H ₂ O) requires :	C,69.9;	H,5.8;	N,7.2%.

Example 14

20

9,10-Dihydro-5-methyl-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide

25

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (0.63g) with Intermediate 2(c) (0.62g) gave, after crystallisation from ethanol, the title compound (0.2g), m.p. 175°.

Analysis Found :	C,71.8;	N,6.2;	N,7.2;
C ₃₄ H ₃₃ N ₃ O ₄ (H ₂ O) requires :	C,72.2;	H,6.2;	N,7.4%.

30

Example 15

35

9,10-Dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-5-methyl-9-oxo-4-acridinecarboxamide

40

A mixture of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.53g) in DMF (30ml) was stirred at room temperature for 10min. Intermediate 16(a) (1.28g) in DMF (20ml) was then added, followed by dicyclohexylcarbodiimide (0.74g) and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The organic layer was then washed with water, dried and concentrated to give a residue which was purified by column chromatography eluting with dichloromethane:methanol (95:5) to give a solid which was recrystallised from ether to give the title compound (0.54g), m.p. 174°.

45

Analysis Found :	C,72.9;	H,6.3;	N,7.4;
C ₃₆ H ₃₇ N ₃ O ₅ requires :	C,73.1;	H,6.3;	N,7.1%.

50

Example 16

55

9,10-Dihydro-5-methoxy-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-9-oxo-4-acridinecarboxamide

A solution of Intermediate 16(a) (1.28g) and dicyclohexylcarbodiimide (0.74g) in DMF (20ml) was added to a stirred solution of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.5g) in DMF (20ml). The resulting mixture was stirred overnight at room temperature, filtered

and concentrated *in vacuo*. The residue was taken up in dichloromethane, and then washed successively with dilute sodium hydroxide solution and water. The organic layer was then dried and evaporated to give a residue which was purified by column chromatography eluting with dichloromethane:methanol (9:1) to give a solid which was crystallised from ether to give the title compound (0.43g), m.p. 188°.

5

Analysis Found :	C,70.9;	H,6.4;	N,7.0;
C ₃₆ H ₃₇ N ₃ O ₆ requires :	C,71.15;	H,6.1;	N,6.9%.

10 The following compounds were prepared in a similar manner to Examples 15 and 16.

Example 17

5-Fluoro-9,10-dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.31g) with Intermediate 8(a) (0.4g) gave, after crystallisation from isopropanol, the title compound (0.2g), m.p. 152°.

20

Analysis Found :	C,85.7;	H,5.6;	F,3.0;	N,6.9;
C ₃₅ H ₃₄ FN ₃ O ₆ (1.5 H ₂ O) requires :	C,85.8;	H,5.8;	F,2.9;	N,6.6%.

Example 18

9,10-Dihydro-5-methoxy-N-[2-methyl-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-9-oxo-4-acridinecarboxamide

30 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.5g) with Intermediate 8(b) (1.3g) gave, after crystallisation from isopropanol/ethanol, the title compound (0.53g), m.p. 160°.

35

Analysis Found :	C,69.6;	H,5.8;	N,6.5;
C ₃₆ H ₃₇ N ₃ O ₆ (0.5H ₂ O) requires :	C,70.1;	H,6.2;	N,6.8%.

Example 19

9,10-Dihydro-5-methyl-N-[2-methyl-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]phenyl]-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 8(b) (1.4g) gave, after crystallisation from acetone, the title compound (0.73g), m.p. 160°.

46

Analysis Found :	C,71.0;	H,6.1;	N,6.5;
C ₃₆ H ₃₇ N ₃ O ₅ (H ₂ O) requires :	C,70.9;	H,6.4;	N,6.9%.

50

Example 20

9,10-Dihydro-5-methoxy-N-[2-methyl-4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-9-oxo-4-acridinecarboxamide

55

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.7g) with Intermediate 16(c) (1.7g) gave, after crystallisation from ethanol, the title compound (0.21g), m.p. 200-201°.

Analysis Found :	C,71.9;	H,5.9;	N,6.9;
C ₃₅ H ₃₅ N ₃ O ₅ (0.5H ₂ O) requires :	C,71.65;	H,6.2;	N,7.2%.

5

Example 215-Fluoro-9,10-dihydro-N-[2-methyl-4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethyl]phenyl]-9-oxo-4-acridinecarboxamide

10

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 16(c) (1.25g) gave, after crystallisation from ethanol, the title compound (0.32g), m.p. 210°.

15

Analysis Found :	C,71.2;	H,5.9;	F,3.4;	N,7.4;
C ₃₄ H ₃₂ FN ₃ O ₄ (0.5 H ₂ O) requires :	C,71.1;	H,5.8;	F,3.3;	N,7.3%.

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Example 229,10-Dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)propoxy]phenyl]-5-methyl-9-oxo-4-acridinecarboxamide

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The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (0.7g) with Intermediate 8(a) (1g) gave, after crystallisation from acetonitrile, the title compound (0.83g), m.p. 183-184°.

Analysis Found :	C,70.2;	H,6.1;	N,6.8;
C ₃₆ H ₃₇ N ₃ O ₆ (0.5H ₂ O) requires :	C,70.1;	H,6.2;	N,6.8%.

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Example 23N-[2-Ethoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)propyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.65g) with Intermediate 16(b) (0.6g) gave, after crystallisation from isopropanol/acetonitrile (9:1), the title compound (0.22g), m.p. 198°.

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Analysis Found :	C,71.1;	H,6.4;	N,6.9;
C ₃₇ H ₃₉ N ₃ O ₆ requires :	C,71.5;	H,6.3;	N,6.8%.

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Example 24N-[2-Methoxy-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propoxy]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

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A mixture of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.5g) in DMF (30 ml) was stirred at room temperature for 10 min. Intermediate 22(b) (1.2g) in DMF (15 ml) was then added, followed by dicyclohexylcarbodiimide (0.8g) and the mixture was stirred at room temperature for 16 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated and the residue was purified by column chromatography eluting with dichloromethane-methanol (97:3). The solid was recrystallised from isopropanol to give the title compound (0.68g). M.p. 108°.

Analysis Found :	C 66.4;	H 5.5;	F 3.0;	N 7.0;
C ₃₄ H ₃₄ FN ₃ O ₅ (H ₂ O) Requires :	C 66.11;	H 5.8;	F 3.1;	N 6.8%.

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Example 25N-[2-Methyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy] phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

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A mixture of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.47g) in DMF (30 ml) was stirred at room temperature for 10 min. Intermediate 22(a) (1.2g) in DMF (15 ml) was then added, followed by dicyclohexylcarbodiimide (0.7g) and the mixture was stirred at room temperature for 16 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated and the residue was purified by column chromatography eluting with dichloromethane-methanol (98:2). The solid was then recrystallised from isopropanol to give the title compound (0.86g). M.p. 130°.

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Analysis Found :	C 69.93;	H 5.89;	F 3.2;	N 7.3;
C ₃₄ H ₃₄ FN ₃ O ₅ Requires :	C 69.97;	H 5.87;	F 3.2;	N 7.2%.

Example 26N-[2-Methoxy-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

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Analysis Found :	C68.4;	H5.9;	N6.7;
C ₃₅ H ₃₇ N ₃ O ₇ Requires :	C68.7;	H6.1;	N6.9%.

In the same way, the following compounds were prepared:

Example 27N-[2-Methyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy] phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridine carboxylic acid (1g) with Intermediate 22(a) (1.23g) gave, after crystallization from isopropanol, the title compound as a solid (1.2g). M.p. 146°.

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N-[2-Methyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.9g) with Intermediate 22(a) (1.2g) gave, after crystallization from isopropanol, the title compound as a solid (1.3g). M.p. 145-150°. NMR includes d 2.2 and 2.3 (2s,2x3H,N-CH₃ and CH₂-Ar), 3.4(s,2H,CH₂-Ar), 3.7(s,6H,OCH₃), 6.6-8.5(m,13H, aromatics).

Example 29

10 N-[2-Methyl-4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethoxy] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridine carboxylic acid (1.2g) with Intermediate 22(d) (1.12g) gave, after crystallization from ethanol, the title compound as a solid (0.6g). M.p. 178-179°.

Analysis Found :	C 70.1;	H 6.1;	N 7.1;
C ₃₄ H ₃₅ N ₃ O ₆ Requires :	C 70.2;	H 6.1;	N 7.2%.

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Example 30

25 N-[2-Ethyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propoxy]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridine carboxylic acid (1g) with Intermediate 22(c) (1.2g) gave, after crystallization from isopropanol, the title compound as a solid (0.95g). M.p. 146°.

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Analysis Found :	C 70.3;	H 6.1;	F 3.2;	N 7.0;
C ₃₅ H ₃₆ FN ₃ O ₅ Requires :	C 70.3;	H 6.1;	F 3.1;	N 7.0%.

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Example 31

N-[2-Methoxy-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propoxy]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

40 The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridine carboxylic acid (0.8g) with Intermediate 22(b) (1.14g) gave, after crystallization from isopropanol, the title compound as a solid (0.4g). M.p. 156-157°.

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Analysis Found :	C 70.6; H 6.3; N 7.15;
C ₃₅ H ₃₇ N ₃ O ₆ Requires :	C 70.6; H 6.3; N 7.05%.

Example 32

50 N-[2-Methyl-4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl] phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridine carboxylic acid (0.82g) with Intermediate 27(a) (1.07g) gave, after crystallization from ethanol, the title compound as a yellow solid (0.21 g). M.p. 125°.

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Analysis Found :	C 68.3;	H 5.8;	F 3.3;	N 7.2;
C ₃₃ H ₃₂ FN ₃ O ₄ (1.5 H ₂ O) Requires :	C 68.3;	H 6.1;	F 3.3;	N 7.2%.

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Example 33

N-[2-Methyl-4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino] ethyl]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

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The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridine carboxylic acid (0.8g) with Intermediate 27(a) (1g) gave, after crystallization from ethanol, the title compound as a yellow solid (0.45g). M.p. 160-161°.

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Analysis Found :	C 73.4;		H 6.3;	N 7.5;
C ₃₄ H ₃₅ N ₃ O ₄ (0.5 H ₂ O) Requires :	C 73.1;	H 6.5;	N 7.5%.	

Example 34

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N-[2-Methoxy-4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino] ethyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

25

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 27(b) (1.3g) gave, after crystallization from ethanol, the title compound as a solid (0.55g). M.p. 161-162°.

Analysis Found :	C69.3;		H5.8;	N7.5;
C ₃₃ H ₃₂ FN ₃ O ₅ Requires :	C69.6;	H5.6;	N7.4%.	

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Example 35

N-[2-Methyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridine carboxylic acid (0.69g) with Intermediate 27(c) (0.65g) gave, after crystallization from isopropanol, the title compound as a solid (0.185g). M.p. 154°.

40

Analysis Found :	C 72.65;		H 6.4;	N 7.0;
C ₃₅ H ₃₇ N ₃ O ₅ Requires :	C 72.5;	H 6.4;	N 7.25%.	

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Example 36

N-[2-Methyl-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

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The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridine carboxylic acid (0.5g) with Intermediate 27(c) (0.59g) gave, after crystallization from isopropanol, the title compound as a solid (0.26 g). M.p. 132°.

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Analysis Found :	C 71.9;	H 6.0;	F 3.3;	N 7.3;
C ₃₄ H ₃₄ FN ₃ O ₄ Requires :	C 71.9;	H 6.0;	F 3.3;	N 7.45%.

Example 37

N-[2-Methoxy-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridine carboxylic acid (0.43g) and Intermediate 30 (0.5g) gave, after crystallization from isopropanol, the title compound as a solid (0.16g). M.p. 105°.

Analysis Found :	C 70.6;	H 6.3;	N 6.9;
C ₃₅ H ₃₇ N ₃ O ₆ Requires :	C 70.6;	H 6.3;	N 7.0%.

10

Example 38N-[2-Methoxy-4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino] propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridine carboxylic acid (0.4g) with Intermediate 30 (0.5g) gave, after crystallization from ethanol/cyclohexane, the title compound as a solid (0.26 g). m.p. 170-190°.

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Analysis Found :	C67.7;	H5.7;	N6.6;
C ₃₄ H ₃₄ FN ₃ O ₅ ,H ₂ O Requires :	C67.9;	H6.0;	N7.0%.

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Example 39N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

30 A mixture of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.42 g) and 1-hydroxybenzotriazole (0.27 g) in DMF (30 ml) was stirred at room temperature for 10 min. Intermediate 33(a) (0.55g) in DMF (30 ml) was then added, followed by dicyclohexylcarbodiimide (0.34 g), and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution, and extracted with dichloromethane. The combined, dried, organic extracts were evaporated to leave an oil which was purified by column chromatography eluting with dichloromethane/methanol (95:5) to give an oil which was crystallised from ethanol and filtered off to give the title compound (0.32g), MP : 131°.

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Analysis Found :	C,71.4;H,5.9;N,7.3;
C ₃₄ H ₃₄ FN ₃ O ₄ Requires :	C,71.9;H,6.0;N,7.4%.

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Example 40N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

50 A mixture of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) and 1-hydroxybenzotriazole (0.41 g) in DMF (50 ml) was stirred at room temperature for 10 min. Intermediate 33(b) (0.9g) in DMF (30 ml) was then added, followed by dicyclohexylcarbodiimide (0.62 g), and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution, and extracted with dichloromethane. The combined, dried, organic extracts were evaporated to leave an oil which was purified by column chromatography eluting with dichloromethane/methanol (95:5) to give a solid. This was crystallised from isopropanol and filtered off to give the title compound (0.31g), MP: 172°.

Analysis Found :	C,71.3;H,6.0;N,7.35;
C ₃₃ H ₃₃ N ₃ O ₅ Requires :	C,71.8;H,6.0;N,7.6%.

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Example 41N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

10 A mixture of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (4 g) and 1-hydroxybenzotriazole (2.83 g) in DMF (50 ml) was stirred at room temperature for 10 min. Intermediate 33(a) (5.5g) in DMF (100 ml) was then added, followed by dicyclohexylcarbodiimide (3.45 g), and the mixture was stirred at room temperature for 16h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution, and extracted with dichloromethane. The combined, dried, organic extracts were evaporated to leave an oil which was purified by column chromatography eluting with dichloromethane/methanol (95:5) to give a solid. This was crystallised from methanol and then filtered off to give the title compound (3.2 g), MP : 140°.

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Analysis Found :	C,74.3;H,6.5;N,7.7;
C ₃₄ H ₃₅ N ₃ O ₄ Requires :	C,74.3;H,6.4;N,7.6%.

Example 42

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N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methyl]amino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

30 A mixture of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) and 1-hydroxybenzotriazole (0.56 g) in DMF (50 ml) was stirred at room temperature for 10 min. Intermediate 33(b) (1g) in DMF (10 ml) was then added followed by dicyclohexylcarbodiimide (0.7 g). The mixture was stirred at room temperature for 16 h and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with dichloromethane. The combined, dried organic extracts were evaporated to leave an oil which was purified by column chromatography eluting with dichloromethane/methanol (9:1) to give a solid. This solid was crystallised from acetonitrile and filtered off to give the title compound (0.35 g), MP : 172°.

Analysis Found :	C,73.6;H,6.0;N,8.0;
C ₃₂ H ₃₁ N ₃ O ₄ Requires :	C,73.7;N,6.0;N,8.1%.

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The following compounds were prepared in a similar manner to Examples 39 to 42 :

Example 43

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N-[4-[[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]thio]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 38(d) (1.16g) gave, after crystallisation from ethanol, the title compound (0.28g), MP : 140°.

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Analysis Found :	C,69.7;H,5.7;N,7.5;
C ₃₃ H ₃₃ N ₃ O ₄ S Requires :	C,69.8;H,5.9;N,7.4 %.

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Example 44N-[4-[2-[(Phenylmethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1 g) with Intermediate 36(c) (1g) gave, after crystallisation from ethanol, the title compound (0.8g), MP : 173°.

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Analysis Found :	C,75.5;H,5.6;N,8.8;
C ₃₀ H ₂₇ N ₃ O ₃ Requires :	C,75.45;H,5.7;N,8.8 %.

Example 45

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N-[4-[3-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

15

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1 g) with Intermediate 38(a) (1.44 g) gave, after crystallisation from ethanol, the title compound (0.82 g), MP : 140°.

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Analysis Found :	C,71.7;H,6.3;N,7.4;
C ₃₄ H ₃₅ N ₃ O ₅ Requires :	C,72.2;H,6.2;N,7.4 %.

Example 46

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N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (2g) with Intermediate 38(c) (2.4g) gave, after crystallisation from isopropanol, the title compound (1.2g), MP : 180°.

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Analysis Found :	C,70.1;	H,6.1;	N,7.2;
C ₃₄ H ₃₅ N ₃ O ₆ requires :	C,70.2;	H,6.1;	N,7.2%.

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Example 47

N-[4-[2-[(2-(4-Methoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

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The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 36(e) (0.9g) gave, after crystallisation from ethanol, the title compound (0.7g), MP : 165°.

45

Analysis Found :	C,73.6;H,6.0;N,8.0;
C ₃₂ H ₃₁ N ₃ O ₄ Requires :	C,73.7;H,6.0;N,8.1%.

Example 48

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N-[4-[3-[(2-(4-Methoxyphenyl)ethyl)methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 38(b) (0.94 g) gave, after crystallisation from ethanol, the title compound (0.9 g), MP : 160°.

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Analysis Found :	C,73.9;H,6.2;N,7.8;
C ₃₃ H ₃₃ N ₃ O ₄ Requires :	C,74.0;H,6.2;N,7.8 %.

Example 49N-[4-[2-[(4-Methoxyphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.6 g) with Intermediate 36(f) (0.72 g) gave, after crystallisation from methanol, the title compound (0.18 g), MP : 146°.

10

Analysis Found :	C,73.5;H,5.8;N,8.1;
C ₃₁ H ₂₉ N ₃ O ₄ Requires :	C,73.35;H,5.8;N,8.3 %.

Example 50N-[4-[2-[(4-Methylphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.7 g) with Intermediate 36(g) (0.78 g) gave, after crystallisation from isopropanol, the title compound (0.23 g), MP : 168°.

20

Analysis Found :	C,75.3;H,6.0;N,8.1;
C ₃₁ H ₂₉ N ₃ O ₃ Requires :	C,75.7;H,5.95;N,8.55 %.

Example 51N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

25 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1 g) with Intermediate 36(b) (1.25 g) gave, after crystallisation from ethanol, the title compound (1.39 g), MP : 140°.

35

Analysis Found :	C,71.7;H,6.2;N,7.7;
C ₃₂ H ₃₁ N ₃ O ₅ Requires :	C,71.5;H,5.8;N,7.8%.

Example 52N-[4-[2-[(4-(Methylthio)phenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

40 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 36(h) (1 g) gave, after crystallisation from ethanol, the title compound (0.75 g), MP : 150°.

45

Analysis Found :	C,71.1;H,5.6;N,7.9;S,5.8; C ₃₁ H ₂₉ N ₃ O ₃ S
Requires :	C,71.1;H,5.6;N,8.0;S,6.1 %.

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Example 53N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-2-(methylthio)-9-oxo-4-acridinecarboxamide

55

The coupling of Intermediate 39 (0.7 g) with Intermediate 36(b) (0.81 g) gave, after crystallisation from ethanol, the title compound (0.45 g), MP : 170°.

Analysis Found :	C,68.1;H,5.65;N,7.0;S,5.4; C ₃₃ H ₃₃ N ₃ O ₅ S
Requires :	C,67.9;H,5.7;N,7.2;S,5.5%.

5

Example 54N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-7-(methylthio)-9-oxo-4-acridinecarboxamide

10

The coupling of 9,10-dihydro-7-(methylthio)-9-oxo-4-acridinecarboxylic acid (0.7 g) with Intermediate 36-(b) (0.81g) gave, after crystallisation from acetonitrile, the title compound (0.14g), MP : 160°.

15

Analysis Found :	C,67.8;H,5.8;N,7.1;S,5.4; C ₃₃ H ₃₃ N ₃ O ₅ S
Requires :	C,67.9;H,5.7;N,7.2;S,5.5 %.

20

Example 55N-[4-[2-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-2-(methylthio)-9-oxo-4-acridinecarboxamide

25

The coupling of Intermediate 39 (0.8g) with Intermediate 36(a) (0.93 g) gave, after crystallisation from ethanol the title compound (0.46g), MP : 150°.

Analysis Found :	C,68.0;H,5.8;N,7.0;S,5.1; C ₃₄ H ₃₅ N ₃ O ₅ S
Requires :	C,68.3;H,5.9;N,7.0;S,5.4 %.

30

Example 56

35

N-[4-[2-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxylic acid (0.72g) with Intermediate 36(a) (0.9g) gave, after crystallisation from isopropanol, the title compound (0.8g), MP : 139°.

40

Analysis Found :	C,72.25;	H,6.2;	N,7.4;
C ₃₄ H ₃₅ N ₃ O ₅ Requires :	C,72.2;	H,6.2;	N,7.4%.

45

Example 57N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

50

The coupling of 9,10-dihydro-5-methoxy-9-oxo-acridinecarboxylic acid (0.8 g) with Intermediate 36(b) (0.94 g) gave, after crystallisation from ethanol, the title compound (0.25 g), MP : 184°.

55

Analysis Found :	C,69.9;H,6.0;N,7.4;
C ₃₃ H ₃₃ N ₃ O ₆ Requires :	C,69.8;H,5.9;N,7.4 %.

Example 58

N-[4-[2-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 36(a) (0.98 g) gave, after crystallisation from ethanol the title compound (0.25 g), MP : 190°.

Analysis Found :	C,70.0;H,6.1;N,7.3;
C ₃₄ H ₃₅ N ₃ O ₅ Requires :	C,70.2;H,6.1;N,7.2 %.

10

Example 59N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1 g) with Intermediate 38(c) (1.4g) gave, after crystallisation from ethanol, the title compound (0.8g), Mp : 130°. IR includes signals at 1650 (CONH), 1620 (CO) and 3350cm⁻¹ (NH).

20

Example 60N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

25

25 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 38(c) (1g) gave, after crystallisation from ethanol, the title compound (0.52g), MP : 150°.

30

Analysis Found :	C,69.6;H,5.7;F,3.25;N,7.3; C ₃₃ H ₃₂ FN ₃ O ₅
Requires :	C,69.6;H,5.7;F,3.3;N,7.4 %.

35

Example 61N-[4-[2-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

40 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.76 g) with Intermediate 33(e) (1g) gave, after crystallisation from acetonitrile, the title compound (0.7g), MP : 180°.

40

Analysis Found :	C,73.5;H,6.1;N,7.9;
C ₃₃ H ₃₃ N ₃ O ₄ Requires :	C,74.0;H,6.2;N,7.8 %.

45

Example 62N-[4-[4-[(4-(Methylthio)phenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

50

50 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(j) (1 g) gave, after crystallisation from acetonitrile, the title compound (0.64g), MP : 135°.

55

Analysis Found :	C,73.7;H,6.2;N,7.9;S,5.7; C ₃₃ H ₃₃ N ₃ O ₂ S
Requires :	C,74.0;H,6.2;N,7.8;S,6.0 %.

Example 63

N-[4-[4-[(4-Fluorophenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.7 g) with Intermediate 33(i) (0.86 g) gave, after crystallisation from acetonitrile, the title compound (0.43 g), MP : 151°.

5

Analysis Found :	C,75.9;H,6.0;F,3.7;N,8.25; C ₃₂ H ₃₀ FN ₃ O ₂
Requires :	C,75.7;H,5.8;F,3.7;N,8.3 %.

10

Example 64N-[4-[3-[(4-Methoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.72 g) with Intermediate 33(g) (0.85 g) gave, after crystallisation from isopropanol, the title compound (0.64 g), MP : 155°.

20

Analysis Found :	C,76.2;H,6.1;N,7.9;
C ₃₂ H ₃₁ N ₃ O ₃ Requires :	C,76.0;H,6.2;N,8.3%.

Example 65N-[4-[4-[(2-(4-Methoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(h) (1 g) gave, after crystallisation from acetonitrile, the title compound (0.53 g), MP : 143°.

30

Analysis Found :	C,76.4;H,6.6;N,7.8;
C ₃₄ H ₃₅ N ₃ O ₃ Requires :	C,76.5;H,6.6;N,7.9 %.

35

Example 66N-[4-[3-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(d) (1 g) gave, after trituration with ether, the title compound (0.88 g), MP : 114°.

45

Analysis Found :	C,74.2;H,6.35;N,7.55;
C ₃₄ H ₃₅ N ₃ O ₄ Requires :	C,74.3;H,6.4;N,7.6 %.

Example 67N-[4-[4-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.72 g) with Intermediate 33(c) (1 g) gave, after crystallisation from acetonitrile, the title compound (0.12 g), MP : 120°.

55

Analysis Found :	C,74.2;H,6.5;N,7.6;
C ₃₅ H ₃₇ N ₃ O ₄ Requires :	C,74.6;H,6.6;N,7.45 %.

Example 68N-[4-[2-[(2-(4-Methoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(k) (0.95 g) gave, after crystallisation from acetonitrile, the title compound (0.4 g), MP : 179°.

10

Analysis Found :	C,76.0;H,6.1;N,8.1;
C ₃₂ H ₃₁ N ₃ O ₃ Requires :	C,76.0;H,6.2;N,8.3 %.

Example 69N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(f) (1 g) gave, after crystallisation from acetonitrile, the title compound (1 g), MP : 112°.

20

Analysis Found :	C,74.1;H,6.2;N,7.7;
C ₃₃ H ₃₃ N ₃ O ₄ Requires :	C,74.0;H,6.2;N,7.8 %.

25

Example 70N-[4-[5-[(3,4-Dimethoxyphenyl)methyl]methylamino]phenyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

30

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(l) (1.15 g) gave, after trituration with ether, the title compound (0.41 g), MP : 110°.

35

Analysis Found :	C,74.3;H,6.6;N,7.4;
C ₃₅ H ₃₇ N ₃ O ₄ Requires :	C,74.6;H,6.6;N,7.45 %.

40

Example 71N-[4-[4-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-7-methoxy-9-oxo-4-acridinecarboxamide

45

The coupling of 9,10-dihydro-7-methoxy-9-oxo-4-acridinecarboxylic acid (1 g) with Intermediate 33(c) (1.3 g) gave, after crystallisation from ethanol, the title compound (0.85 g), MP : 155°.

50

Analysis Found :	C,72.7;H,6.9;N,7.05;
C ₃₆ H ₃₉ N ₃ O ₅ Requires :	C,72.8;H,6.6;N,7.1 %.

Example 72N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

55 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(a) (0.98 g) gave, after crystallisation from isopropanol, the title compound (0.12 g), MP : 157°.

Analysis Found :	C,71.9;H,6.4;N,7.2;
Requires :	C,72.5;H,6.4;N,7.25 %.

5

Example 73N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

10

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.72 g) with Intermediate 33(f) (0.9g) gave, after crystallisation from ethanol, the title compound (0.89 g), MP : 158°.

15

Analysis Found :	C,71.9;H,6.1;F,3.25;N,7.7; C ₃₃ H ₃₂ FN ₃ O ₄
Requires :	C,71.65;H,5.8;F,3.4;N,7.6 %.

20

Example 74N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

25

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 33(b) (1.2 g) gave, after crystallisation from ethanol, the title compound (0.78 g), MP : 175°.

Analysis Found :	C,69.9;H,5.5;F,3.1;N,7.45; C ₃₂ H ₃₀ FN ₃ O ₄
(0.5 H ₂ O) Requires :	C,70.1;H,5.7;F,3.5;N,7.65%.

30

Example 75N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxylic acid (0.6g) with Intermediate 33(a) (0.7 g) gave, after crystallisation from acetonitrile, the title compound (0.35 g), MP : 174°.

40

Analysis Found :	C,68.6;H,5.7;N,9.5;
C ₃₄ H ₃₄ N ₄ O ₆ Requires :	C,68.7;H,5.8;N,9.4 %.

45

Example 76N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxamide

50

The coupling of 9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxylic acid (0.6 g) with Intermediate 33(b) (0.63 g) gave, after crystallisation from isopropanol, the title compound (0.45 g), MP : 197°.

55

Analysis Found :	C,67.4;H,5.3;N,9.7;
C ₃₂ H ₃₀ N ₄ O ₆ Requires :	C,67.8;H,5.3;N,9.9 %.

Example 77

N-[4-[5-[(3,4-Dimethoxyphenyl)methyl]methylamino]pentyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(l) (1 g) gave, after crystallisation from acetonitrile, the title compound (0.29 g); MP : 130°.

Analysis Found :	C,71.8;H,6.2;F,3.2;N,7.1; C ₃₅ H ₃₆ FN ₃ O ₄
Requires :	C,72.3;H,6.2;F,3.3;N,7.2 %.

10

Example 78N-[4-[4-[(2-(4-Methoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(h) (0.93 g) gave, after trituration with ether, the title compound (0.31 g), MP : 182°.

20

Analysis Found :	C,74.2;H,6.6;N,7.8;
C ₃₅ H ₃₇ N ₃ O ₄ Requires :	C,74.6;H,6.6;N,7.5 %.

Example 79N-[4-[2-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

30 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(e) (0.94 g) gave, after crystallisation from isopropanol, the title compound (0.17 g), MP : 179°.

35

Analysis Found :	C,72.3;H,6.0;N,7.8;
C ₃₄ H ₃₅ H ₃ O ₅ Requires :	C,72.2;H,6.2;N,7.4 %.

Example 80N-[4-[4-[(2-(3,4-Dimethoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

45 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(c) (1 g) gave, after crystallisation from isopropanol, the title compound (0.12 g), MP : 170°. IR gave signals at 1645 (CONH), 1620 (CO) and 3300cm⁻¹ (NH).

Example 81N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxamide

50 The coupling of 9,10-dihydro-5-nitro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(f) (0.88 g) gave, after crystallisation from isopropanol, the title compound (0.29 g), MP : 192°.

55

Analysis Found :	C,67.8;H,5.6;N,9.4;
C ₃₃ H ₃₂ N ₄ O ₆ Requires :	C,68.3;H,5.6;N,9.65 %.

Example 82

N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(f) (0.93 g) gave, after crystallisation from ethanol, the title compound (0.27 g), MP : 180°.

10

Analysis Found :	C,72.0;H,6.1;N,7.6;
C ₃₄ H ₄₅ N ₃ O ₅ Requires :	C,72.2;H,6.2;N,7.4 %.

Example 83

N-[4-[2-[(Phenylmethyl)ethylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 36(i) (0.9 g) gave, after crystallisation from ethanol, the title compound (0.34 g), MP : 157°.

20

Analysis Found :	C,75.3;H,5.9;N,8.4;
C ₃₁ H ₂₉ N ₃ O ₃ Requires :	C,75.7;H,5.9;N,8.5 %.

25

Example 84

N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxamide

30 The coupling of 9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxylic acid (0.8 g) with Intermediate 33(a) (1.04 g) gave, after crystallisation from isopropanol, the title compound (0.65 g), MP : 142°. IR gave signals at 1675 (CONH), 1610 (CO) and 3250cm⁻¹ (NH).

35

Example 85

N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxamide

40

The coupling of 9,10-dihydro-10-methyl-9-oxo-4-acridinecarboxylic acid (0.87 g) with Intermediate 33(h) (1g) gave, after crystallisation from isopropanol, the title compound (0.42 g), MP : 182°.

45

Analysis Found :	C,73.5;H,6.1;N,7.8;
C ₃₃ H ₃₃ N ₃ O ₄ Requires :	C,74.0;H,6.2;N,7.8 %.

Example 86

50

N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-7-methoxy-9-oxo-4-acridinecarboxamide

55 The coupling of 9,10-dihydro-7-methoxy-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 33(a) (0.97g) gave, after crystallisation from isopropanol, the title compound (0.17g), MP : 172°.

Analysis Found :	C,71.5;	H,6.4;	N,6.9;
C ₃₅ H ₃₇ N ₃ O ₅ , 0.5H ₂ O Requires :	C,71.4;	H,6.5;	N,7.1%.

5

Example 87

N-[4-[[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]thio] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

10

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.7g) with Intermediate 36(d) (1g) gave, after crystallisation from isopropanol, the title compound (0.26g), MP : 113°.

15

Analysis Found :	C,69.3;	H,5.5;	N,7.4;	S,5.8;
C ₃₂ H ₃₁ N ₃ O ₄ S Requires :	C,69.4;	H,5.6;	N,7.6;	S,5.8%.

Example 88

20

N-[4-[[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

25

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 38(d) (1.09g) gave, after crystallisation from ethanol, the title compound (50mg), MP : 158°.

Analysis Found :	C,69.4;	H,5.9;	N,6.9;	S,5.6;
C ₃₄ H ₃₅ N ₃ O ₄ S, 0.5 H ₂ O Requires :	C,69.1;	H,6.1;	N,7.1;	S,5.4%.

30

Example 89

35

N-[4-[[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 38(d) (1.28g) gave, after crystallisation from acetonitrile, the title compound (0.37g), MP : 184-186°.

40

Analysis Found :	C,68.1;	H,5.9;	N,6.8;	S,5.2;
C ₃₄ H ₃₅ N ₃ O ₅ S Requires :	C,68.3;	H,5.9;	N,7.0;	S,5.4%.

45

Example 90

N-[4-[[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-fluoro-9-oxo-4-acridinecarboxamide

50

The coupling of 9,10-dihydro-5-fluoro-9-oxo-acridinecarboxylic acid (0.9g) with Intermediate 38(d) (1.1g) gave, after crystallisation from isopropanol, the title compound (0.5g), MP : 120-130°.

55

Analysis Found :	C,66.6;	H,5.6;	F,3.1;	N,6.9;	S,5.3;
C ₃₃ H ₃₂ FN ₃ O ₄ S, 0.5 H ₂ O Requires :	C,66.6;	H,5.6;	F,3.2;	N,7.1;	S,5.4%.

Example 91

N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methylthio-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-5-methylthio-9-oxo-4-acridinecarboxylic acid (0.7g) with Intermediate 33(b) (0.74g) gave, after crystallisation from ethanol, the title compound (0.3g), MP : 190°.

Analysis Found :	C,68.5;	H,6.1;	N,7.2;
C ₃₃ H ₃₃ N ₃ O ₄ S, 0.5 H ₂ O Requires :	C,68.7;	H,5.9;	N,7.3%.

10

Example 92N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1.27g) with Intermediate 33(b) (1.5g) gave, after crystallisation from isopropanol/diisopropylether, the title compound (0.3g), MP : 119°.

20

Analysis Found :	C,73.5;	H,6.2;	N,7.6;
C ₃₃ H ₃₃ N ₃ O ₄ Requires :	C,74.0;	H,6.2;	N,7.8%.

25

Example 93N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

30

The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 38(c) (1.3g) gave, after crystallisation from isopropanol, the title compound (0.9g), MP : 160°.

35

Analysis Found :	C,72.3;	H,6.3;	N,7.5;
C ₃₄ H ₃₅ N ₃ O ₅ requires :	C,72.2;	H,6.3;	N,7.5%.

Example 94

40

N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethylamino] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.4g) with Intermediate 43 (1.4g) gave after crystallisation from isopropanol, the title compound (0.2g), MP : 196°.

45

Analysis Found :	C,69.8;	H,6.3;	N,10.0;
C ₃₃ H ₃₄ N ₄ O ₅ requires :	C,69.9;	H,6.1;	N,9.9%.

50

Example 95N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5,8-dimethoxy-9-oxo-4-acridinecarboxamide

55

The coupling of 9,10-dihydro-5,8-dimethoxy-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 33-(b) (0.67g) gave, after crystallisation from ethanol, the title compound (0.15g) MP : 196°.

Analysis Found :	C,68.99;	H,5.76;	N,7.18;
C ₃₄ H ₃₅ N ₃ O ₆ , 0.5 H ₂ O Requires :	C,69.13;	H,6.14;	N,7.11%.

5

Example 96N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5,7-dimethoxy-9-oxo-4-acridinecarboxamide

10

The coupling of Intermediate 44 (1.4g) with Intermediate 33(b) (1.2g) gave, after crystallisation from ethanol, the title compound (0.25g), MP > 260°.

15

Analysis Found :	C,70.09;	H,6.35;	N,7.01;
C ₃₄ H ₃₅ N ₃ O ₆ Requires	C,70.20;	H,6.06;	N,7.22%.

Example 97

20

N-[4-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-6,7,8-trimethoxy-9-oxo-4-acridinecarboxamide

25

The coupling of Intermediate 45 (0.6g) with Intermediate 33(b) (0.6g) gave, after crystallisation from isopropanol, the title compound (0.4g), MP : 158°.

Analysis Found :	C,68.69;	H,6.32;	N,6.40;
C ₃₅ H ₃₇ N ₃ O ₇ Requires	C,68.72;	H,6.10;	N,6.87%.

30

Example 98

35

N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]amino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

40

A mixture of Intermediate 40 (0.5g) and 3,4-dimethoxybenzenemethanamine (0.5 g) was heated for 1 h at 140°. Water was then added and the mixture was extracted with dichloromethane. The dried organic phase was concentrated to give a solid which was purified by column chromatography eluting with dichloromethane/methanol (9:1). The resulting solid was crystallised from benzene to give the title compound (50 mg), MP : 138-139°.

Analysis Found :	C,70.1;H,5.9;N,7.5;
C ₃₂ H ₃₁ N ₃ O ₅ (0.5 H ₂ O) Requires	C,70.3;H,5.9;N,7.7%.

45

Example 99

50

Oxalate of N-[4-[[(3,4-Dimethoxyphenyl)methyl]methylamino] butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

A solution of Example 41 (0.55 g) and oxalic acid dihydrate (0.126 g) in ethanol (10 ml) was boiled for 2 min. After cooling and scratching, crystallisation took place. The crystals were filtered off and dried to afford the title compound (0.55 g), MP : 155-160°.

55

Analysis Found :	C,66.3;H,5.9;N,6.3;
C ₃₆ H ₃₇ N ₃ O ₂ (0.5 H ₂ O) Requires :	C,66.6;H,5.9;N,6.4%.

5

Example 100Maleate of N-[4-[4-[(3,4-dimethoxyphenyl)methyl] methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridin-ecarboxamide

10

A solution of Example 41 (0.55 g) and maleic acid (0.130 g) in ethanol (50 ml) was boiled for 2 min. After cooling and scratching, crystallisation took place. The crystals were filtered off and dried to afford the title compound (0.5 g), MP : 205°.

15

Analysis Found :	C,68.2;H,5.9;N,6.2;
C ₃₈ H ₃₉ N ₃ O ₈ Requires :	C,68.5;H,5.9;N,6.3%.

20

Example 101Hydrochloride of N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino] butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

25

A hot solution of Example 41(0.55 g) in ethanol (50 ml) was treated with a slight excess of an ethereal solution of hydrochloric acid. The solution was then concentrated to give a foam which was triturated with isopropanol to afford the title compound (0.4 g) as crystals, MP : 165°.

30

Analysis Found :	C,67.6;H,6.3;N,7.0;
C ₃₄ H ₃₆ ClN ₃ O ₄ . H ₂ O Requires :	C,67.5;H,6.4;N,7.0%.

Example 102

35

L+ lactate of N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino] butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

40

A solution of Example 41 (0.55 g) and L+ lactic acid (0.95 g) in isopropanol (30 ml) was boiled for 2 min. After cooling and scratching, crystallisation took place. The crystals were filtered off and dried to afford the title compound (0.45 g), MP : 120°.

45

Analysis Found :	C,69.5;H,6.5;N,6.6;
C ₃₇ H ₄₁ N ₃ O ₇ Requires :	C,69.4;H,6.6;N,6.5%.

Example 103

50

Oxalate of N-[3-[(3,4-dimethoxyphenyl)methyl] methylamino]propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

55

A mixture of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) and 1-hydroxybenzotriazole (0.63g) in DMF (30ml) was stirred at room temperature for 10 min. Intermediate 51 (1.23g) in DMF (3.9ml) was then added followed by dicyclohexylcarbodiimide (0.8g) and the mixture was stirred at room temperature for 16 hours and then filtered. The filtrate was concentrated in vacuo, treated with dilute sodium hydroxide solution and extracted with methylene chloride. The combined, dried, organic extracts were evaporated to leave an oil which, after purified by column chromatography on silica gel eluting with

methylene chloride/methanol (99:1), led to the title compound (1.1g), m.p. 126°.

5	Analysis Found : $C_{33}H_{32}F_1N_3O_4.C_2H_2O_4$ (H ₂ O) Requires :	C,63.9; C,63.5;	H,5.4; H,5.5;	F,2.8; F,2.9;	N,6.2; N,6.3%
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The following compounds were prepared in a similar manner to Example 103:

Example 104

10 N-[3-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.5g) with Intermediate 48(b) (1.22g) gave, after crystallisation from isopropanol, the title compound (0.47g) as a solid, m.p. 124°.

20	Analysis Found : $C_{34}H_{35}N_3O_6$ Requires :	C,70.1; C,70.2;	H,6.1; H,6.1;	N,7.05; N,7.2%
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Example 105

25 Oxalate of N-[3-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

30 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridine carboxylic acid (1.26g) with Intermediate 51 (1.23g) gave the title compound (1.13g), m.p. 112-114°.

35	Analysis Found : $C_{34}H_{35}N_3O_5.C_2H_2O_4$ (0.5 H ₂ O) Requires :	C,65.2; C,65.0;	H,6.2; H,5.8;	N,6.2; N,6.3%
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36 Example 106

Fumarate of N-[3-[2-[(3,4-dimethoxyphenyl)methyl]methylamino] ethyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

40 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.34g) with Intermediate 48(a) (0.4g) gave the title compound (0.3g), m.p. 155°.

Example 107

45 Fumarate of N-[3-[2-[(3,4-dimethoxyphenyl)methyl]methylamino] ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

50 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.36g) with Intermediate 48(a) (0.4g) gave the title compound (0.13g), m.p. 140°.

55 Example 108

N-[4-[4-[(3,4-Dimethoxyphenyl)methyl]methylamino]butyl]-2-methoxyphenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

55 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.38g) with Intermediate 55 (0.5g) gave, after crystallisation from isopropanol, the title compound (0.36g) as a solid, MP : 114-115°.

Analysis Found :	C,70.98;	H,6.19;	N,6.79;	C ₃₆ H ₃₉ N ₃ O ₆
Requires :	C,70.92;	H,6.45;	N,6.89%.	

5

Example 109

9,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethyl]amino]-phenyl]-4-acridinecarboxamide

10

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.99g) with Intermediate 59 (1.2g) gave, after crystallisation from acetonitrile, the title compound (1.3g), MP : 228 - 234°.

15

Analysis Found :	C,69.27;	H,5.87;	N,9.37;
C ₃₄ H ₃₄ N ₄ O ₅ , 0.5H ₂ O Requires :	C,69.48;	H,6.00;	N,9.50%.

20

Example 110

N-[4-[2-(2,3-Dihydro-5,6-dimethoxy-1H-isoindol-2-yl)ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

25

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.54g) with Intermediate 60 (0.6g) gave after crystallisation from ethanol, the title compound (0.3g), MP : 215 - 225°. NMR includes signals at d 2.85(4H,s,N-(CH₂)₂-Ph); 3.7(6H,s,2xOMe); 3.8(3H,s,OMe); 3.9(4H,s,2xN-CH₂-Ph).

35

Example 111

30

9,10-Dihydro-5,8-dimethoxy-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)propyl]-phenyl]-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5,8-dimethoxy-9-oxo-4-acridinecarboxylic acid (0.7g) with Intermediate 16-(a) (0.83g) gave, after crystallisation from ethanol, the title compound (0.1g), MP : 140°.

Analysis Found :	C,67.44;	H,5.94;	N,6.80;
C ₃₇ H ₃₉ N ₃ O ₇ , H ₂ O Requires :	C,67.77;	H,6.30;	N,6.40%.

40

Example 112

45

9,10-Dihydro-5-methoxy-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)-1-hydroxyethyl]phenyl]-9-oxo-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.49g) with Intermediate 63 (0.5g) gave, after crystallisation from acetonitrile, the title compound (0.8g), MP : 160-165°.

50

Analysis Found :	C,68.51;	H,5.74;	N,7.25;
C ₃₄ H ₃₃ N ₃ O ₆ , H ₂ O Requires :	C,68.33;	H,5.90;	N,7.09%.

55

Example 113

9,10-Dihydro-5-methoxy-9-oxo-N-[4-[[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethyl]-methylamino]methyl]phenyl]-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.53g) Intermediate 67 (0.7g) gave, by precipitation from methylene chloride/diethyl ether, the title compound (0.5g), MP : 202°.

5	Analysis Found : C ₃₆ H ₃₈ N ₄ O ₅ , 1.25H ₂ O Requires :	C,68.68; C,68.71;	H,6.27; H,6.48;	N,8.52; N,8.90%.
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Example 114

10 N-[4-[[2-[(3,4-Dimethoxyphenyl)methyl]methylamino]ethyl] methylamino]methyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.1g) with Intermediate 70 (1.43g) gave, after crystallisation from methanol, the title compound (0.75g) as yellow crystals, MP : 170°.

20	Analysis Found : C ₃₅ H ₃₈ N ₄ O ₅ , 0.5 H ₂ O Requires :	C,69.69; C,69.63;	H,6.30; H,6.51;	N,9.10; N,9.28%.
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Example 115

25 5-Fluoro-9,10-dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)propyl]phenyl]-9-oxo-4-acridinecarboxamide

30 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.5g) with Intermediate 16(a) (0.63g) gave, after crystallisation from ethanol, the title compound (0.3g), MP : 128°. NMR includes signals at d 3.6(3H,s,OMe); 3.8(6H,s,2xOMe); 9.15(1H,s,NH^{CO}); 11.35(1H,s,NH acridone).

Example 116

35 N-[4-[[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-(methylthio)-9-oxo-4-acridinecarboxamide

40 The coupling of 9,10-dihydro-5-methylthio-9-oxo-4-acridinecarboxylic acid (0.3g) with Intermediate 38(d) (0.36g) gave, after crystallisation from methanol, the title compound (0.13g), MP : 142°. NMR includes signals at d 2.2(3H,s,SMe); 2.45(3H,s,NMe); 3.7(6H,s,2xOMe).

Example 117

45 N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]propyl]-2-methoxyphenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide

50 The coupling of 9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxylic acid (0.75g) and Intermediate 30 (1g) gave, after crystallisation from methanol, the title compound (0.1g), MP : 111°. NMR includes signals at d 2.18(3H,s,NCH₃); 2.55(3H,s,CH₃ acridone); 3.42(2H,s,N-CH₂-Ph); 3.9(9H,3s,3xOMe).

Example 118

55 N-[2-Ethoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl) propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

55 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 16(b) (0.86g) gave, after crystallisation from acetonitrile, the title compound (0.4g), MP : 200°. NMR includes signals at d 1,4(2H,t,CH₃-CH₂); 3,7(6H,s,2xOMe).

Example 119

N-[4-[[(3,4-Dimethoxyphenyl)methyl]methylamino]-2-butenyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (154mg) with Intermediate 72 (210mg) gave, after crystallisation from ethanol, the title compound (80mg), MP : 140°.

Analysis Found :	C,74.17;	H,6.08;	N,7.61;
C ₃₄ H ₃₃ N ₃ O ₄ Requires :	C,74.55;	H,6.07;	N,7.67%.

10

Example 120N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]-1-propenyl] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.95g) with Intermediate 74 (1.1g) gave, after crystallisation from ethanol, the title compound (0.7g), MP : 200°.

20

Analysis Found :	C,72.46;	H,6.04;	N,7.61;
C ₃₄ H ₃₃ N ₃ O ₅ Requires :	C,72.45;	H,5.90;	N,7.45%.

25

Example 1215-Methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6-methoxy-2-isoquinoliny)ethyl]phenyl]-9,10-dihydro-4-acridinecarboxamide

30 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.5g) with Intermediate 76 (0.48g) gave, after crystallisation from pyridine/water, the title compound (0.4g), MP: 260°.

35

Analysis Found :	C,74.29;H,6.06;N,8.02; C ₃₃ H ₃₁ N ₃ O ₄
requires :	C,74.28;H,5.88;N,7.87%

Example 1225-Fluoro-9,10-dihydro-9-oxo-N-[3-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-4-acridinecarboxamide

40 The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 79 (1.3g) gave, after crystallisation from isopropanol, the title compound (0.25g), MP: 128°.

45

Analysis Found :	C,68.84;H,5.67;F,3.01;N,6.88;
C ₃₄ H ₃₂ FN ₃ O ₄ (1.5H ₂ O) requires :	C,68.90;H,5.95;F,3.20;N,7.09%

50

Example 1239,10-Dihydro-5-methoxy-9-oxo-N-[3-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-4-acridinecarboxamide

55

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.2g) with Intermediate 79 (1.2g) gave, after crystallisation from isopropanol, the title compound (0.5g), MP: 138-140°.

Analysis Found :	C,70.55;H,6.25;N,7.06;
C ₃₅ H ₃₅ N ₃ O ₅ (H ₂ O) requires :	C,70.56;H,6.26;N,7.05%

5

Example 124

N-[4-[3-[(3,4-Dimethoxyphenyl)methyl]methylamino]-2-hydroxypropoxy] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide

10

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 81 (1.3g) gave, after crystallisation from isopropanol, the title compound (0.7g), MP: 175°.

15

Analysis Found :	C,68.38;H,5.82;N,6.86; C ₃₄ H ₃₅ N ₃ O ₇
requires :	C,68.33;H,5.90;N,7.03%

Example 125

20

9,10-Dihydro-5-methoxy-9-oxo-N-[4-[3-[(3,4,5-trimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-4-acridinecarboxamide

25

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (1.5g) with Intermediate 83 (1.3g) gave, after crystallisation from isopropanol, the title compound (1.3g), MP: 186°.

Analysis Found :	C,68.82;H,6.08;N,6.83; C ₃₅ H ₃₇ N ₃ O ₇
requires :	C,68.72;H,6.10;N,6.87%

30

Example 126

35

Fumarate of 5-fluoro-9,10-dihydro-N-[2-methoxy-5-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (1g) with Intermediate 86 (1.2g) gave the title compound (0.5g), MP: 166-168°.

40

Analysis Found :	C,63.78;H,5.15;N,6.10;
C ₃₈ H ₃₆ FN ₃ O ₉ (H ₂ O) requires :	C,63.76;H,5.35;N,5.87%

45

Example 127

9,10-Dihydro-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-2-isoquinoliny)propoxy]phenyl]-4-acridinecarboxamide

50

The coupling of 9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 88 (0.9g) gave, after crystallisation from ethanol, the title compound (0.3g), MP: 182°.

Analysis Found :	C,74.88;H,5.81;N,8.16;
C ₃₂ H ₂₉ N ₃ O ₃ (0.5H ₂ O) requires :	C,74.98;H,5.90;N,8.20%

55

Example 128

9,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-7-methoxy-2-isoquinolinyl)ethyl]phenyl]-4-acridinecarboxamide

5 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.7g) with Intermediate 90 (0.7g) gave, after crystallisation from isopropanol, the title compound (0.65g), MP: 213-216°.

Analysis Found :	C,73.27;H,5.94;N,7.82;
C ₃₃ H ₃₁ N ₃ O ₄ (0.5H ₂ O) requires :	C,73.04;H,5.94;N,7.74%

10

Example 129

15 9,10-Dihydro-5-methoxy-9-oxo-N-[3-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethyl]phenyl]-4-acridinecarboxamide

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.5g) with Intermediate 92 (0.57g) gave, after crystallisation from isopropanol, the title compound (0.15g), MP: 152°.

20

Analysis Found :	C,71.33;H,5.77;N,7.16;
C ₃₄ H ₃₃ N ₃ O ₅ (0.5H ₂ O) requires :	C,71.30;H,5.98;N,7.33%

25

Example 130

5-Fluoro-9,10-dihydro-9-oxo-N-[3-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinolinyl)ethyl]phenyl]-4-acridinecarboxamide

30

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.5g) with Intermediate 92 (0.57g) gave, after crystallisation from isopropanol, the title compound (0.35g), MP: 178°.

35

Analysis Found :	C,70.80;H,5.36;F,3.34;N,7.34;
C ₃₃ H ₃₀ FN ₃ O ₄ (0.5H ₂ O) requires :	C,70.70;H,5.57;F,3.38;N,7.49%

Example 131

40 Fumarate of N-[5-[2-[(3,4-Dimethoxyphenyl)methyl]methylamino] ethyl]-2-methoxyphenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide

The coupling of 5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxylic acid (0.8g) with Intermediate 95 (1g) gave the title compound (0.5g), MP: 140-142°.

45

Analysis Found :	C,62.4;H,5.1;N,5.8;
C ₃₇ H ₃₆ FN ₃ O ₉ (1.5H ₂ O) requires :	C,62.35;H,5.5;N,5.9%

50

Example 132

9,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-5,6-dimethoxy-2-isoquinolinyl)ethyl]phenyl]-4-acridinecarboxamide

55

The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.19g) with Intermediate 97 (0.22g) gave, after crystallisation from pyridine/water, the title compound (0.32g). MP:235-237°. NMR includes signals at d 2.6-3.0 (8H,m,2x N-(CH₂)₂-Ar), 3.6 (2H,s,N-CH₂-Ar), 3.75 (6H,bs,OCH₃), 4 (3H,s,OCH₃),

6.5-8.5 (12H,m,aromatics).

5

Analysis Found :	C,72.38;H,5.80;N,7.41;
C ₃₄ H ₃₃ N ₃ O ₅ requires :	C,72.45;H,5.90;N,7.45%.

Example 133

10 9,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7,8-trimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide

15 The coupling of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (0.26g) with Intermediate 99 (0.3g) gave, after crystallisation from isopropanol, the title compound (0.3g). MP:222-226°. NMR includes signals at d 2.4-2.9 (8H,m,2x N-(CH₂)₂-Ar), 3.45 (2H,s,N-CH₂-Ar), 3.7 (9H,bs,OCH₃), 3.9 (3H,s,OCH₃), 6.2-8.4 (11H,m,aromatics).

20

Analysis Found :	C,69.46;	H,6.14;	N,6.84;
C ₃₅ H ₃₅ N ₃ O ₆ (0.5 H ₂ O) requires:	C,69.75;	H,6.02;	N,6.97%.

Example 134

25 5-Amino-N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide

30 A suspension of Example 75 (0.15g) in ethanol (40ml) was hydrogenated at room temperature in presence of 10% palladium-on- carbon (70mg). After the hydrogen absorption was completed, the mixture was diluted with methylene chloride (50ml). The catalyst was filtered off and the solution concentrated in vacuo to give the title compound (85mg) as a yellow solid, MP : 250°.

35

Analysis Found :	C,72.38;	H,6.69;	N,9.06;
C ₃₄ H ₃₆ N ₄ O ₄ Requires :	C,72.31;	H,6.42;	N,9.92%.

Example 135

40 9,10-Dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide

45 Dicyclohexylcarbodiimide (22.76g) in DMF (50ml) was added dropwise to a stirred mixture of 9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxylic acid (28.9g) and 1-hydroxybenzotriazole hydrate (15.66g) in DMF (300ml) maintained at 0°, followed by Intermediate 101 (33.5g) in DMF (150ml). After 4 hours at 0° and 2 days at room temperature, the mixture was filtered, the filtrate was concentrated in vacuo and the residue taken up in 1N sodium hydroxide and extracted with dichloromethane. The organic layer was then washed with water, dried and evaporated to give a solid residue. This was dissolved in 500ml of boiling pyridine and the solution was clarified by filtration. The clear solution was diluted with 10ml of water and the product crystallised on cooling to give the title compound (52.82g). M.p. : 215-225°. NMR includes d 2.60-2.95 (m,8H,CH₂); 3.58 (s,2H,N-CH₂-Ph); 3.72 (s,6H,OMe); 4.05 (s,3H,OMe acridone); 6.78 (2s,2H,Anisoquinoline), 7.20-7.88 (m,8H,Ar.), 8.48 (t,2H,H₁ and H₈ acridone), 10.60 (s,1H,CONH), 12.32 (s,1H,NH acridone).

55

Analysis found :	C,72.07;	H,5.96;	N,7.35;
C ₃₄ H ₃₃ N ₃ O ₅ requires :	C,72.45;	H,5.90;	N,7.45%.

Example 136

Maleate salt of 9,10-dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)-ethyl]phenyl]-4-acridinecarboxamide

5

Example 135 (100mg) was dissolved in 50ml of a mixture of dichloromethane and methanol (1:1) and maleic acid (22mg) was added. The mixture was boiled until a clear solution was obtained and the solution was evaporated in vacuo. The residue was taken up in hot methanol and cooled to give the title compound as yellow needles (90mg). M.P.: 171 to 187°.

10

In the same way the following salts of Example 135 were prepared:

15

Fumarate :	m.p. : 170-203°.
Succinate :	m.p.: 135-143°.
L (+) Tartrate :	m.p.: 165-180°.

Example 137

20

Hydrochloride salt of 9,10-dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)-ethyl]phenyl]-4-acridinecarboxamide

25

Example 135 (100mg) was dissolved in a mixture of methanol and dichloromethane (4:1) and excess methanolic hydrogen chloride was added. The solvate was recovered which after addition of diethyl ether and filtration gave the title compound (ca. 100mg). MP 225° (softens with progressive loss of solvent).

30

Example 138

In vitro cytotoxicity of MDR inhibitors in Chinese Hamster Ovary cells

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The multidrug resistant Chinese Hamster Ovary (CHO) cell line CH^RC5 was obtained from Dr V Ling, Princess Margaret Hospital, Toronto, Canada and maintained as anchorage-dependent monolayers in a minimum essential medium supplemented with thymidine, adenosine, 10% fetal bovine serum, 2mM L-glutamine (Flow), 100 units/ml penicillin and 100mg/ml streptomycin in a humidified atmosphere of 95% air and 5% carbon dioxide. Cells were passaged into culture flasks twice a week after dissociation with EDTA.

40

CH^RC5 cells were seeded at a density of 10⁴ cells/well in microtitre plates. After 24 hours, the medium was removed and replaced by 0.1ml of fresh medium containing successive two-fold dilutions of MDR inhibitors. Each MDR inhibitor was assayed in duplicate in two-fold dilution from 1250 to 20nM. The last well of each column was utilised to verify the lack of toxicity at the top dose of the MDR inhibitor in the absence of doxorubicin. Other control conditions were assayed on each microtitre plate : cells alone (1 well), doxorubicin alone (7 wells), amiodarone (a range of two-fold dilutions starting at 5mM; two wells each). 0.1ml of a 10mg/ml solution of doxorubicin was added. After 72 hours incubation cell viability was assessed by the reduction of 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT; Sigma) to a dark blue formazan product. In particular, 20ml of a 5mg/ml solution of MTT prepared in phosphate buffered saline was added to each well. After 4 hours incubation at 37°, the medium was aspirated and replaced with 0.1ml dimethylsulphoxide. After vigorous shaking, the quantity of formazan product formed was assessed by its optical density at 550nm. The absorbance is directly related to the number of surviving cells in the wells.

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Cytotoxicity calculations were performed on the average of the two wells for each condition. The concentration of each MDR inhibitor giving a 50% reduction of the optical density relative to cells treated with doxorubicin alone was determined to give an EC₅₀ value.

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Results

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In the above test the compounds of the specific Examples hereinabove had EC₅₀ values in the range of 0.018 to 0.72mM. Thus, for example, the compound of Example 1 had an EC₅₀ of 0.02mM, at least 100 times more potent than prototype MDR inhibitors including amiodarone (EC₅₀ 3mM) and verapamil (3mM).

Administration of the compound of Example 1 to mice orally produced no visible toxic effects at single doses up to 300mg/kg.

The following are examples of pharmaceutical compositions according to the invention. The term 'Active Ingredient' as used hereinafter means a compound of the invention and may be for example the compound of Example 1.

5 Example A - Oral Tablet

		Per Tablet (mg)
10	Active Ingredient	50.0
	Microcrystalline Cellulose	110.0
	Lactose	67.5
	Sodium Starch Glycolate	20.0
	Magnesium Stearate	2.5
15	Total	250.0

The drug is sieved through a 250mm sieve and then the five powders are intimately mixed in a blender and compressed on 3/8 inch standard concave punches in a tabletting machine.

20 Example B - Oral Capsule

		Per Capsule (mg)
25	Active Ingredient	50.0
	Microcrystalline Cellulose	66.5
	Lactose USP	66.5
	Sodium Starch Glycolate	15.0
30	Magnesium Stearate	2.0
	Total	200.0

The drug is sieved through a 250mm sieve and then the five powders are intimately mixed in a blender and filled into No. 2 hard gelatin capsule shells on a capsule filling machine.

35

Example C - Injection for Intravenous Administration (10mg in 10mL)

40

% w/w

45	Active Ingredient	0.1
	Cancer chemotherapy agent	as required
	Water for Injection to	100.0
50	Dilute hydrochloric acid to	pH 3.0

The active ingredient (and cancer chemotherapy agent where appropriate) is dissolved with mixing in the Water For Injection, adding acid slowly until the pH is 3.0. The solution is sparged with nitrogen and filtratively sterilized through a sterilized filter of 0.22 micron pore size. Under aseptic conditions this sterile solution is placed into sterile ampoules and the ampoules flame sealed.

Example D - Oral Syrup

5

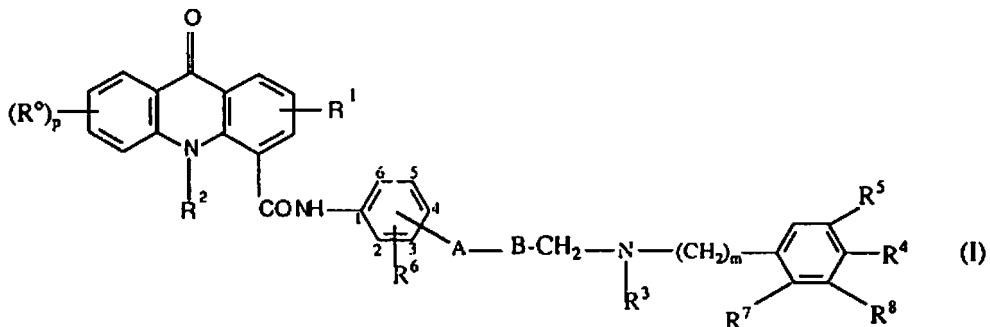
% w/v

	Active Ingredient	2.0
10	Cancer chemotherapy agent	as required
	Dilute hydrochloric acid to	pH 3.0
15	Sorbitol solution	60 v/v
	Flavour	as required
	Distilled water to	100

20 The active ingredient (and cancer chemotherapy agent where appropriate) is dissolved in some of the water with stirring by adding gradually the hydrochloric acid until the pH is 3.0. The sorbitol solution, flavour and the rest of the water are added and the pH re-adjusted to 3.0. The syrup is clarified by filtration through suitable filter pads.

25 **Claims**

1. A compound of formula (I)



wherein R⁰ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio, amino or nitro group;

p represents 1; or when R⁰ represents C₁₋₄alkoxy may also represent 2 or 3;

R¹ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy or C₁₋₄alkylthio group;

R² represents a hydrogen atom or a C₁₋₄alkyl group;

A represents an oxygen or a sulphur atom, a bond or a group (CH₂)₁NR³ (where 1 represents zero or 1 and R³ represents a hydrogen atom or a methyl group);

B represents a C₁₋₄ alkylene chain optionally substituted by a hydroxyl group, except that the hydroxyl group and moiety A cannot be attached to the same carbon atom when A represents an oxygen or sulphur atom or a group (CH₂)₁NR³, or when A represents a bond B may also represent a C₂₋₄alkenylene chain;

R³ represents a hydrogen atom or a C₁₋₄alkyl group;

m represents 1 or 2;

R⁴ represents a hydrogen or a halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy or C₁₋₄alkylthio group;

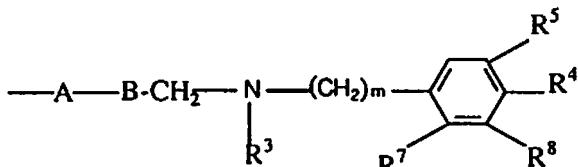
R⁵ represents a hydrogen atom or a C₁₋₄alkoxy group;

R⁶ represents a hydrogen atom or a C₁₋₄alkyl or C₁₋₄alkoxy group;

R⁷ represents a hydrogen atom or R³ and R⁷ together form a group -(CH₂)_n-where n represents 1 or 2;

R⁸ represents a hydrogen atom or a C₁₋₄alkoxy group;
the group

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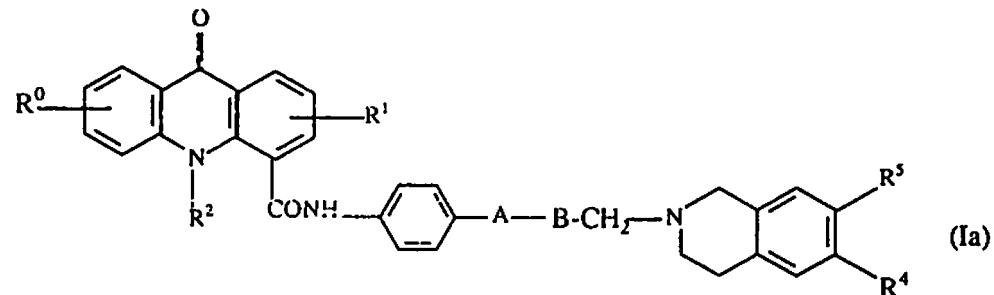
is attached at the benzene ring 3 or 4 position relate to the carboxamide substituent, provided that when the group is attached at the benzene ring 3 position then R⁶ must be attached at the benzene ring 6 position;

15 and salts and solvates thereof.

2. A compound according to Claim 1 in which R⁹ represents a hydrogen or fluorine atom, or a C₁₋₄alkoxy, C₁₋₄alkyl or C₁₋₄alkylthio group and R¹ represents a hydrogen atom.
- 20 3. A compound according to Claim 1 or Claim 2 in which an R⁹ group is situated at the 5-position of the acridone molecule.
4. A compound according to any preceding claim in which R² represents a hydrogen atom.
- 25 5. A compound according to any preceding claim in which R⁴ and R⁵ each represent a C₁₋₄alkoxy group and R⁸ represents a hydrogen atom.
6. A compound according to any preceding claim in which m represents 1 and R³ and R⁷ together form a group -(CH₂)₂-.
- 30 7. A compound of formula (Ia)

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wherein R⁰ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylthio or nitro group;

R¹ represents a hydrogen or halogen atom, or a C₁₋₄alkyl, C₁₋₄alkoxy or C₁₋₄alkylthio group;

R² represents a hydrogen atom or a C₁₋₄alkyl group;

A represents an oxygen or a sulphur atom or a bond;

50 B represents an unsubstituted C₁₋₄alkylene chain;

R⁴ and R⁵ each independently represents a C₁₋₄alkoxy group; and physiologically acceptable salts and solvates thereof.

8. A compound according to Claim 7 in which R⁹ represents a hydrogen or fluorine atom or a C₁₋₄alkoxy or C₁₋₄alkyl group, R¹ and R² each represent a hydrogen atom and R⁴ and R⁵ each represent a C₁₋₄alkoxy group.

- 55 9. A compound according to Claim 8 in which the R⁹ group is situated at the 5-position of the acridone

molecule.

10. A compound according to Claim 1 which is 9,10-dihydro-5-methoxy-9-oxo-N-[4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)ethyl]phenyl]-4-acridinecarboxamide and physiologically acceptable salts and solvates thereof.

11. A compound according to Claim 1 selected from:
 9,10-dihydro-5-methoxy-9-oxo-N-[4-[[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]-phenyl]-4-acridinecarboxamide;
 5-fluoro-9,10-dihydro-9-oxo-N-[4-[[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]-phenyl]-4-acridinecarboxamide;
 9,10-dihydro-5-methoxy-9-oxo-N-[4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propoxy]-phenyl]-4-acridinecarboxamide;
 9,10-dihydro-5-methyl-9-oxo-N-[4-[[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]thio]-phenyl]-4-acridinecarboxamide;
 9,10-dihydro-5-methoxy-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]-phenyl]-9-oxo-4-acridinecarboxamide;
 9,10-dihydro-N-[2-methoxy-4-[3-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isoquinoliny)propyl]phenyl]-5-methyl-9-oxo-4-acridinecarboxamide;
 and physiologically acceptable salts and solvates thereof.

12. A compound according to Claim 1 selected from :
 N-[4-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-4-[(3,4-dimethoxyphenyl)methyl]methylamino]butyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-3-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-5-[(3,4-dimethoxyphenyl)methyl]methylamino]pentyl]phenyl]-5-fluoro-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-3-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-[(3,4-dimethoxyphenyl)methyl]methylamino]propyl]thio] phenyl]-9,10-dihydro-5-fluoro-9-oxo-4-acridinecarboxamide;

N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methylthio-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(3,4-dimethoxyphenyl)methyl]methylamino]ethyl]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide;
 5 N-[4-[3-[(3,4-dimethoxyphenyl)methyl]methylamino]propoxy]phenyl]-9,10-dihydro-5-methyl-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 10 N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]butyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(2-(4-methoxyphenyl)ethyl)methylamino]ethyl]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-2-(methylthio)-9-oxo-4-acridinecarboxamide;
 15 N-[4-[3-[(2-(3,4-dimethoxyphenyl)ethyl)methylamino]propoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(2-(4-methoxyphenyl)ethyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(3,4-dimethoxyphenyl)methyl)methylamino]ethoxy]phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 20 N-[4-[3-[(3,4-dimethoxyphenyl)methyl)methylamino]propoxy]phenyl]-9,10-dihydro-5-methoxy-9-oxo-4-acridinecarboxamide;
 N-[4-[2-[(3,4-dimethoxyphenyl)methyl)methylamino]ethyl]thio)phenyl]-9,10-dihydro-9-oxo-4-acridinecarboxamide;
 25 and physiologically acceptable salts and solvates thereof.

13. A compound according to any preceding claim for use in therapy.
14. A compound according to any preceding claim for use in the treatment of a mammal which is suffering from cancer, to improve or increase the efficacy of an antitumour drug, or increase or restore sensitivity of a tumour to an antitumour drug, or reverse or reduce resistance of a tumour to an antitumour drug.
15. Use of a compound according to any of Claims 1 to 12 for the manufacture of a medicament for the treatment of a mammal suffering from cancer, to improve or increase the efficacy of an antitumour drug, or increase or restore sensitivity of a tumour to an antitumour drug, or reverse or reduce resistance of a tumour to an antitumour drug.
16. A method of treatment of a mammal which is suffering from cancer, which method comprises administering to said mammal an effective amount of a compound according to any of Claims 1 to 12 to improve or increase the efficacy of an antitumour drug, or increase or restore sensitivity of a tumour to an antitumour drug, or reverse or reduce resistance of a tumour to an antitumour drug.
17. A pharmaceutical composition which comprises a compound according to any of Claims 1 to 12 together with one or more physiologically acceptable carriers or excipients.
18. A pharmaceutical composition which comprises an active amount of a compound according to any of Claims 1 to 12 for use in the treatment of a mammal which is suffering from cancer, to improve or increase the efficacy of an antitumour drug, or increase or restore sensitivity of a tumour to an antitumour drug, or reverse or reduce resistance of a tumour to an antitumour drug.
19. A pharmaceutical composition according to Claim 17 or Claim 18 comprising a compound according to Claim 10.
20. A pharmaceutical composition according to any of Claims 17 to 19 in a form suitable for oral, buccal, parenteral or rectal administration.
21. A pharmaceutical composition according to any of Claims 17 to 20 in unit dosage form.

22. A product containing a compound according to any of Claims 1 to 12 and an antitumour drug as a combined preparation for simultaneous, separate or sequential use in treating cancer.

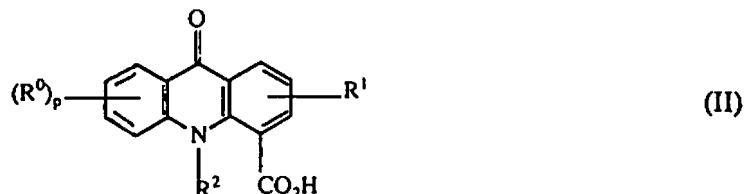
23. A compound according to any of Claims 1 to 12 and an antitumour drug in the presence of each other in the human or non-human animal body for use in treating cancer.

24. Product or process according to any of Claims 14 to 23 (except Claim 17) wherein the antitumour drug is selected from Vinca alkaloids, anthracyclines, taxol and derivatives thereof, podophyllotoxins, mitoxantrone, actinomycin, colchicine, gramicidine D, amsacrine or any drug having cross-resistance with the above drugs characterised by the so-called MDR phenotype.

25. A process for the preparation of a compound according to Claim 1 which comprises:

(A) reacting a compound of formula (II)

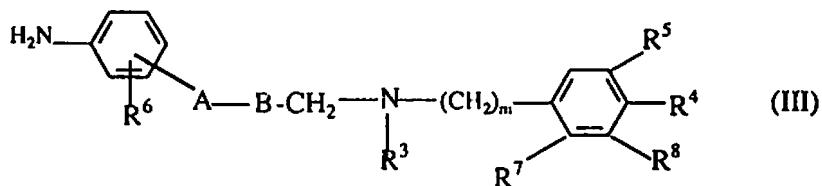
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with a compound of formula (III)

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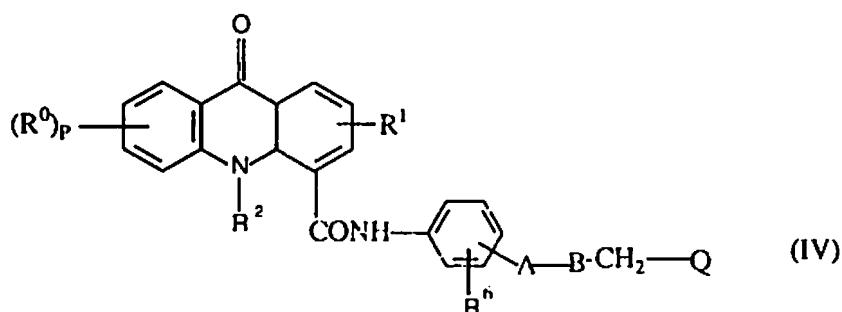


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in the presence of a coupling reagent; or
 (B) reacting a compound of formula (IV)

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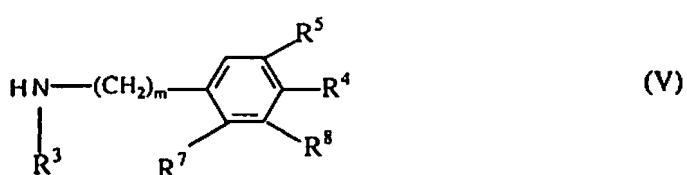


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(wherein Q represents a halogen atom) with a compound of formula (V)

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or a salt thereof in the presence of an acid acceptor; with salt formation as an optional step subsequent

to process (A) or (B).

26. Compounds according to any of Claims 1 to 12 substantially as herein described.

5 27. Compositions according to any of Claims 17 to 21 substantially as herein described.

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PARTIAL EUROPEAN SEARCH REPORT

which under Rule 45 of the European Patent Convention
shall be considered, for the purposes of subsequent
proceedings, as the European search report

Application Number

EP 92 10 0123

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	EP-A-0 098 098 (DEVELOPMENT FINANCE CORP. OF NEW ZEALAND) * Claims * ---	1,13-23	C 07 D 219/06 C 07 D 401/12 A 61 K 31/435 A 61 K 31/47
A	JOURNAL OF MEDICINAL CHEMISTRY, vol. 31, no. 3, March 1988, pages 707-712, Washington, US; B.D. PALMER et al.: "Potential antitumor agents. 54. Chromophore requirements for in vivo antitumor activity among the general class of linear tricyclic carboxamides" * The whole document * ---	1,13-23	
A	JOURNAL OF MEDICINAL CHEMISTRY, vol. 30, no. 4, April 1987, pages 658-663, Washington, US; W.A. DENNY et al.: "Potential antitumor agents. 49. 5-Substituted derivatives of N-[2-(Dimethylamino)ethyl]-9-aminoacridine-4-carboxamide with in vivo solid-tumor activity" * The whole document * -----	1,13-23	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 D
INCOMPLETE SEARCH			
<p>The Search Division considers that the present European patent application does not comply with the provisions of the European Patent Convention to such an extent that it is not possible to carry out a meaningful search into the state of the art on the basis of some of the claims</p> <p>Claims searched completely :</p> <p>Claims searched incompletely :</p> <p>Claims not searched :</p> <p>Reason for the limitation of the search:</p> <p>Remark: Although claims 16,22,23 are directed to a method of treatment of the human/animal body (Art. 52(4) EPC) the search has been carried out and based on the alleged effects of the compound/composition</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	05-03-1992	HENRY J.C.	
CATEGORY OF CITED DOCUMENTS		<p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- & : member of the same patent family, corresponding document</p>	
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p>			